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(54) Title: TRANSLATIONAL PROFILING

(57) Abstract: Polypeptides representative of proteins expressed by a given cell type and isolated nucleic acids that encode the polypeptides are disclosed. The compositions and method described can be used to define a cell type at a given developmental, metabolic, or disease stage by identifying and cataloging proteins expressed in the cell. The compositions can also be used in the manufacture of therapeutics as well as in diagnostics and drug screening.

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TRANSLATIONAL PROFILING

CROSS REFERENCE TO RELATED APPLICATIONS

This application claims priority from U.S. Provisional Application No.

- 5 60/279,495, filed March 28, 2001, U.S. Provisional Application No. 60/292,544, filed
May 21, 2001, U.S. Provisional Application No. 60/310,801, filed August 8, 2001, U.S.
Provisional Application No. 60/326,370, filed October 1, 2001, U.S. Provisional
Application No. 60/336,780, filed December 4, 2001, and U.S. Provisional Application
No. 60/358,985, filed February 20, 2002. These applications are incorporated herein by
10 reference in their entirety.

FIELD OF THE INVENTION

The invention relates to peptides identified by translational profiling methods, as
well as nucleic acids encoding the peptides, methods of using the peptides to characterize
15 the protein composition of a cell, and methods of using the peptides to diagnose, prevent,
and treat disease.

REFERENCE TO SEQUENCE LISTING SUBMITTED ON A COMPACT DISC

This application includes a compact disc (four copies of disc submitted)
20 containing a sequence listing. The sequence listing is identified on the compact disc as
follows.

File Name	Date of Creation	Size (bytes)
08191-026WO1.TXT	March 25, 2002	8,015,000

The entire content of the sequence listing is herein incorporated by reference.

25

BACKGROUND OF THE INVENTION

Essentially every cell within an organism contains the complete and identical
genetic information of that organism, but expresses only a subset of that total
complement of genes. For example, the human genome, which is composed of a total of
30 three billion nucleotides, is currently thought to include approximately 30,000-40,000

genes. However, individual cells expresses only about 2,000 to about 4,000 different proteins, corresponding to only 10% of the total number of genes. It is the concerted activity of the proteins expressed in a given cell that orchestrates the activities that define a particular cell type at a given developmental, metabolic or disease stage.

5 In the past decades it has become clear that the development and the pathology of many diseases involves differences in gene expression. Indeed, healthy and diseased tissue or cell types can frequently be distinguished by differences in gene expression. For example, normal cells may evolve to highly invasive and metastatic cancer cells by activation of certain growth-inducing genes, e.g., oncogenes, or the inactivation of certain
10 growth-inhibitory genes, e.g., tumor suppressors or apoptosis activators. Levine, 1997, *Cell* 88:323; Hunter, 1997, *Cell* 88:333; Jacobson, 1997, *Cell* 88:347; Nagata, 1997, *Cell* 88:355; Fraser *et al.*, 1996, *Cell* 85:781. Altered expression of such genes, e.g., growth activators or growth suppressors, in turn affects expression of other genes. See, The National Cancer Institute, "The Nation's Investment In Cancer Research: A Budget
15 Proposal For Fiscal Years 1997/98", Prepared by the Director, National Cancer Institute, pp. 55-77.

Pathological gene expression differences are not confined to cancer. Autoimmune disorders, many neurodegenerative diseases, inflammatory diseases, restenosis, atherosclerosis, many metabolic diseases, and numerous other disorders are
20 believed to involve aberrant expression of particular genes. Naparstek *et al.*, 1993, *Ann. Rev. Immunol.* 11:79; Sercarz *et al.*, 1993, *Ann. Rev. Immunol.* 11:729. As a consequence, a challenge in medical research is to understand the role each gene or its encoded protein plays in maintaining normal cellular homeostasis and to utilize this heightened understanding in improving the ability to treat disease and/or identify
25 predispositions to disease at stages when treatment and/or prevention methods are available.

Significant resources have been expended to identify and isolate genes relevant to disease development. One approach has been to sequence and catalogue all the individual genes contained in the genome of a species. In the case of humans, the NIH
30 initiated the Humane Genome Project in 1990, with the goal to sequence the entire human genome by the year 2005. Stephens *et al.*, 1990, *Science* 250:237; Cantor, 1990,

Science 248:49-51. The near complete sequence of the human genome was published in advance of the 2005 target date. Venter et al., *Science* 2001 291:1304; International Human Genome Sequencing Consortium *Nature* 2001 409:860. However, the vast amount of information made available by the sequencing of the human genome is 5 insufficient to resolve the mysteries of many disease processes because cellular function and dysfunction results from the concerted interaction and differential expression of proteins. Indeed, nucleotide sequence information alone does not indicate when, where, and how much of a given gene is expressed at the protein level.

10

SUMMARY OF THE INVENTION

The present invention is based on the purification of a series of peptide sequences derived from proteins produced within a panel of cells. The purification and sequencing of these peptides demonstrates both the existence of a given protein as well as the production of the given protein in a particular cell type. In many cases, the existence of a 15 given protein was uncertain prior to the characterization described herein, as it had never previously been isolated or even detected. Members of one class of peptides described herein, termed expressed protein tags (EPTs), bind to and are presented by human MHC class I or class II molecules. Members of a second class of peptides are chemically or enzymatically prepared from complex protein mixtures.

20

The invention generally relates to novel peptides and proteins containing the novel amino acid sequences. In addition, the invention relates to nucleic acids encoding polypeptides containing the novel peptides, methods of using the peptide sequences in the context of a database or a peptide profile to characterize the protein composition of a cell or a peptide array comprising peptides of the invention, and using the identified peptides 25 and corresponding nucleic acids in methods of treatment, diagnosis, and screening.

In one aspect, the invention features a purified polypeptide including a peptide sequence selected from the group consisting of SEQ ID NOs:1-235. In an embodiment, the polypeptide comprises at least 8 contiguous amino acids of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235. In another embodiment, the invention 30 features a purified immunogenic polypeptide comprising at least 8 contiguous amino acids of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

"Immunogenic peptides" are peptides that result in or enhance an immune response in a mammal. Examples of immunogenic peptides can be found, for example in U.S. 5,827,516 and U.S. 6,183,746. In another embodiment, the invention features a purified polypeptide, comprising at least an immunogenic portion of a protein, wherein the protein comprises an 5 amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

In another aspect, the invention features a purified polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235, wherein the purified polypeptide comprises at least 25 amino acids. In an example, the purified polypeptide comprises fewer than 100 amino acids. In another example, the purified 10 polypeptide comprises fewer than 50 amino acids.

In one embodiment, the polypeptide consists of a peptide sequence selected from the group consisting of SEQ ID NOs: 1-235. In another embodiment, the polypeptide consists essentially of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

15 The peptide sequence can be identical to that of a naturally processed class I MHC-binding peptide. Alternatively, the peptide sequence can be identical to that of a naturally processed class II MHC-binding peptide.

In another aspect, the invention features an isolated nucleic acid encoding a polypeptide comprising a peptide sequence selected from the group consisting of SEQ ID 20 NOs: 1-235. In an embodiment, the polypeptide comprises an amino acid sequence which is at least 95% identical to an amino acid selected from the group consisting of SEQ ID NOs: 1-235. In another embodiment, the isolated nucleic acid comprises a nucleotide sequence that encodes a polypeptide comprising an amino acid sequence selected from the group consisting of a variant of any one of SEQ ID NOs: 1-235, wherein the variant has no more 25 than two conservative amino acid substitutions. In a further embodiment, the isolated nucleic acid comprises a nucleotide sequence that encodes a polypeptide comprising at least 8 contiguous amino acids of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

30 In some examples the encoded polypeptide includes a peptide sequence identical to that of a naturally processed class I MHC-binding peptide. Alternatively, the peptide sequence can be identical to that of a naturally processed class II MHC-binding peptide.

- In one embodiment, an isolated nucleic acid encodes a polypeptide including a peptide sequence identical to a segment of a naturally occurring protein, wherein the peptide sequence is selected from the group consisting of SEQ ID NOs: 1-235, and wherein the polypeptide does not include more than 10, 15, 20, 30, 40, 50, 60, 70 , 80 , 90, or 100 consecutive amino acids identical to a portion of the naturally occurring protein. The peptide sequence can be identical to that of a naturally processed class I MHC-binding peptide. Alternatively, the peptide sequence can be identical to that of a naturally processed class II MHC-binding peptide.

5 In an aspect, the invention features an isolated nucleic acid comprising a nucleotide sequence encoding a polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235. In another aspect, the isolated nucleic acid comprises a nucleotide sequence encoding a polypeptide consisting essentially of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

10 The invention also includes an expression vector containing a nucleic acid described herein. In an example, the vector comprises expression control sequences that direct expression of the polypeptide. In another example, the vector comprises expression control sequences that direct expression of the nucleic acid molecule. Also included in the invention is a cell containing an expression vector of the invention.

15 In another aspect, the invention features an antibody specific for a polypeptide of the invention, e.g., a peptide sequence selected from the group consisting of SEQ ID NOs: 1-235. In an example, the antibody selectively binds to the polypeptide which is expressed on a cell surface. In another example, the antibody of the polypeptide is a target of a second antibody located on a cell surface.

20 In another aspect, the invention features a humanized antibody which specifically binds to a domain of a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235 or an isolated nucleic acid which encodes the antibody. In preferred embodiments, the humanized antibody is a full length antibody, a human IgG, an antibody fragment and a F(ab)₂. The invention also features a humanized antibody as described herein bound to a detectable label. In another aspect, the invention 25 30 features an immobilized antibody comprising a humanized antibody as described herein

bound to a solid phase. In a further aspect, the invention features a conjugate comprising a humanized antibody as described herein bound to a cytotoxic agent.

The invention also includes a method for determining the presence of a protein comprising exposing a sample suspected of containing the protein to a humanized antibody as described herein and determining binding of the antibody to the sample. In another aspect, the invention includes a kit comprising a humanized antibody as described herein and instructions for using the humanized antibody to detect a protein that binds to the antibody.

The invention also includes a method of making an antibody, the method comprising: (a) providing a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235 or a nucleic acid encoding such a polypeptide to a mammal in an amount effective to induce the production of an antibody that binds to the polypeptide; (b) isolating from the mammal a cell that produces an antibody that selectively binds to a polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235; (c) immortalizing the cell isolated in step (b); and (d) isolating antibodies from the immortalized cell.

The invention also includes a method of modulating the activity of a polypeptide described herein, the method including contacting the polypeptide with a compound that binds to the polypeptide in a concentration sufficient to modulate the activity of the polypeptide. In an example, the compound that binds the polypeptide is an antibody that selectively binds a polypeptide consisting of an amino acid sequence selected for the group consisting of SEQ ID NOs: 1-235.

In another aspect, the invention features a method of treating a disorder in a mammal, the method including: (1) identifying a mammal with the disorder; and (2) administering to the mammal a compound that modulates the expression or activity of a polypeptide described herein, wherein the administration results in an amelioration of one or more symptoms of the disorder. The disorder can be for example a cellular proliferative and/or differentiative disorder or a disorder associated with the particular biological class of proteins to which the polypeptide belongs.

30 . . . In another aspect, the invention features a method for detecting the presence of a polypeptide described herein in a sample, the method including: (1) contacting the

sample with a compound that selectively binds to the polypeptide; and (2) determining whether the compound binds to the polypeptide in the sample.

In another aspect, the invention features a method for detecting the presence of a disorder in a mammal, the method including: (1) providing a biological sample derived from the mammal; (2) contacting the sample with a compound that binds to a polypeptide described herein or to a nucleic acid that encodes such a polypeptide; and (3) determining whether the compound binds to the sample, wherein binding of the compound to the sample indicates the presence or absence of the disorder in the mammal.

In another aspect, the invention features a method for imaging a site in a mammal, the method including: (1) administering a compound to a mammal, wherein the compound binds to a polypeptide described herein (or to a nucleic acid that encodes such a polypeptide) at the site in the mammal; and (2) detecting the compound with an imaging detector, to thereby image the site in the mammal.

In another aspect, the invention features a method for identifying a compound that modulates the activity of a polypeptide described herein, the method including: (1) contacting a polypeptide described herein with a test compound; and (2) determining the effect of the test compound on the activity of the polypeptide, to thereby identify a compound that modulates the activity of the polypeptide.

In another aspect, the invention features a method for identifying a compound that modulates the expression of a nucleic acid described herein, the method including: (1) contacting the nucleic acid with a test compound; and (2) determining the effect of the test compound on the expression of the nucleic acid, to thereby identify a compound that modulates the expression of the nucleic acid.

In another aspect, the invention features a peptide profile that is characteristic for a given cell, wherein the profile includes a representation of at least ten different polypeptides in the cell, wherein each of the at least ten different polypeptides contains a peptide selected from the group consisting of SEQ ID NOs: 1-235, and wherein the peptide profile is a reproducible characteristic of the cell. In one example, the each of the at least ten different polypeptides contains an MHC-binding peptide. In one example, the representation characterizes each individual peptide based upon at least one physical or chemical attribute, the at least one physical or chemical attribute including amino acid

sequence. In addition, the representation can characterize each individual peptide based upon at least two physical or chemical attributes, e.g., wherein one of the physical or chemical attributes is amino acid sequence. For example, one of the physical or chemical attributes can be mass-to-charge ratio or ion-fragmentation pattern. In another example, the 5 representation can characterize each individual peptide based upon at least three physical or chemical attributes. In another aspect, the invention features a polypeptide profile that is characteristic of a selected cell under selected conditions, wherein the profile comprises a representation of at least ten different polypeptides expressed by the cell, wherein each of the at least ten different polypeptides comprises an amino acid sequence selected from the 10 group consisting of SEQ ID NOs: 1-235, and wherein the polypeptide profile is a reproducible characteristic of the cell.

In another aspect, the invention features a database, stored on a machine-readable medium, containing: two categories of data respectively representing (a) peptide profiles and (b) cell sources; and associations among instances of the two categories of data, wherein 15 the data representing peptide profiles include a peptide profile described herein, and wherein the database configures a computer to enable finding instances of data of one of the categories based on their associations with instances of data the other category.

In another aspect, the invention features a database, stored on a machine-readable medium, comprising: (a) three categories of data respectively representing (i) 20 polypeptides, (ii) cell sources, and (iii) cell treatments; and (b) associations among instances of the three categories of data, wherein the data representing peptides comprises at least 100 polypeptides each having an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235, and wherein the database configures a computer to enable finding instances of data of one of the categories based on their associations with 25 instances of data of at least one other category.

In another aspect, the invention features a peptide array comprising at least 100 peptides selected from the group consisting of peptides consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235, each peptide linked to a solid support at a known location. In another aspect the invention features a 30 collection of at least 10 polypeptide arrays, each array comprising at least 100

polypeptides consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235, each peptide linked to a solid support at a known location.

In another aspect, the invention features a method of selecting an antibody, the method including: (1) contacting a polypeptide described herein with an *in vitro* library
5 of antibodies; (2) binding an antibody to the polypeptide; and (3) selecting the antibody that binds to the polypeptide.

In another aspect, the invention features an immunogenic composition comprising a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235, the composition when injected into a mammal elicits an immunogenic
10 response directed against a polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

The invention also features a method for treating a cancer comprising administering to a patient in need of such treatment an amount of a composition comprising a polypeptide comprising an amino acid sequence selected from the group
15 consisting of SEQ ID NOs: 1-235 in an amount sufficient to elicit an immunogenic response.

The invention also features a method for treating a cancer patient, the method comprising administering to the patient an antibody that selectively binds to a peptide consisting of an amino acid sequence selected from the group consisting of SEQ ID
20 NOs:1-235.

In another aspect, the invention features a method for identifying a compound that binds to a naturally processed class I or class II MHC-binding polypeptide, the method comprising exposing a test compound to a collection of at least 100 polypeptides selected from the group consisting of polypeptides having an amino acid sequence selected from
25 the group consisting of SEQ ID NOs:1-235, and identifying a peptide to which the test compound binds.

An "isolated" or "purified" polypeptide, protein, or peptide (these terms are used interchangeably) is a polypeptide, protein, or peptide that is separated from those components (proteins and other naturally-occurring organic molecules) that naturally
30 accompany it. Typically, the polypeptide, protein, or peptide is substantially pure when it constitutes at least 60%, by weight, of the protein in the preparation. Preferably, the

protein in the preparation consists of at least 75%, more preferably at least 90%, and most preferably at least 99%, by weight, of the polypeptide, protein, or peptide of the invention.

- An "isolated" or "purified" nucleic acid refers to a nucleic acid that is separated from other nucleic acid molecules present in the natural source of the nucleic acid. With regards to genomic DNA, the term "isolated" refers to a nucleic acid molecule that is free of sequences that naturally flank the nucleic acid (i.e., sequences located at the 5' and/or 3' ends of the nucleic acid) in the genomic DNA of the organism from which the nucleic acid is derived. For example, in various embodiments, the isolated nucleic acid molecule can contain less than about 5 kb, 4 kb, 3 kb, 2 kb, 1 kb, 0.5 kb or 0.1 kb of 5' and/or 3' nucleotide sequences that naturally flank the nucleic acid molecule in genomic DNA of the cell from which the nucleic acid is derived. Moreover, an "isolated" nucleic acid molecule, such as a cDNA molecule, can be substantially free of other cellular material, or culture medium when produced by recombinant techniques, or substantially free of chemical precursors or other chemicals when chemically synthesized.

- The term "nucleic acid" includes, for example, a recombinant DNA that is incorporated into a vector such as an autonomously replicating plasmid or virus. The nucleic acids herein can comprise ribonucleotides, deoxyribonucleotides, or modified forms of either nucleotide. Isolated nucleic acid sequences can be single or double stranded and can be polynucleotides or oligonucleotides.

- Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Suitable methods and materials are described below, although methods and materials similar or equivalent to those described herein can also be used in the practice or testing of the present invention. All publications, patent applications, patents, and other references mentioned herein are incorporated by reference in their entirety. In case of conflict, the present specification, including definitions, will control. In addition, the materials, methods, and examples are illustrative only and not intended to be limiting.

- Other features and advantages of the invention will be apparent from the following detailed description, and from the claims.

DETAILED DESCRIPTION

The present invention relates generally to peptide sequences identified by translational profiling methods. The invention also relates to polypeptides containing the peptide sequences, nucleic acids encoding polypeptides containing the peptide sequences, the use of these compositions in methods and systems for analyzing the protein composition of cells and cell populations, and methods of using the compositions in the diagnosis and treatment of disease as well as in the screening for therapeutic compounds to treat disease.

10

Polypeptides and Nucleic Acids

The invention features purified polypeptides comprising a peptide sequence of any of SEQ ID NOS: 1-235. Polypeptides can be purified from cells or tissue sources using a variety of protein purification techniques.

15

Methods of obtaining a purified preparation of a recombinant protein are well known in the art and include culturing transformed host cells under culture conditions suitable to express the protein, and purifying the resulting protein using known purification processes, such as gel filtration or ion exchange chromatography. The purification of the protein may also utilize an affinity column containing agents which will bind to the protein; one or more column steps over affinity resins such as concanavalin A-agarose, heparin-toyopearl® or Cibacrom blue 3GA Sepharose®, one or more steps involving hydrophobic interaction chromatography using such resins as phenyl ether, butyl ether, or propyl ether; and/or immunoaffinity chromatography.

20

Additionally, one or more reverse-phase high performance liquid chromatography (RP-HPLC) steps employing hydrophobic RP-HPLC media, e.g., silica gel having pendant methyl or other aliphatic groups, can be employed to further purify the protein. Some or all of the foregoing purification steps, in various combinations, can also be employed to provide a substantially homogenous isolated protein. The protein thus purified is substantially free of other mammalian proteins and is defined in accordance with the present invention as a "purified polypeptide."

A polypeptide can also be isolated from cells or tissue sources by using an affinity molecule to separate the polypeptide from a complex mixture of proteins. For example, a polypeptide can be purified by isolating a molecule, e.g., an MHC class I or class II molecule, to which the polypeptide is bound and eluting the polypeptide from the

5 molecule. Alternatively, a polypeptide can be isolated from cells or tissue sources by using an anti-polypeptide antibody, e.g., an antibody described herein. Polypeptides or fragments thereof can also be synthesized chemically, e.g., by solid phase methods using an automated peptide synthesizer. Polypeptides can also be isolated and fragmented *in vitro* by the action of chemical or enzymatic treatments.

10 The amino acid sequences of the peptides of SEQ ID NOs: 1-235 are presented in Table 1 (see Examples). This table indicates the "source protein symbols" from which each of the peptides is derived. Symbols are obtained from three places in the following order: (a) gene symbol(s) and alias(es) from Locus Link; (b) gene name(s) from LocusLink; or (c) Locus titles from LocusLink. The table also provides SEQ ID NOs for
15 each of the source proteins. The sequences corresponding to the SEQ ID NOs of these source proteins were obtained from GenBank™ accession numbers. The accession numbers can be viewed by entering (under a "Protein" search) the sequence for the "source protein reference" at www.ncbi.nlm.nih.gov/PubMed/. The entire content of each of this references is herein incorporated by reference. Many of the respective
20 GenBank™ accessions also provide a reference to a nucleic acid sequence encoding the source protein. These nucleic acid sequences are also incorporated by reference in their entirety.

In some embodiments, the polypeptide does not include more than 200 consecutive amino acids, e.g., no more than 150, 100, 90, 80, 70, 60, 50, 40, or 30 amino
25 acids, identical to a portion of a naturally occurring protein from which a peptide of SEQ ID NOs: 1-235 is derived. In other embodiments, the polypeptide consists of a peptide of any of SEQ ID NOs: 1-235, or a variant peptide as described below. In other embodiments, the polypeptide comprises at least 8 contiguous amino acids of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235. In another
30 embodiment, the purified polypeptide comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235, wherein the purified polypeptide comprises

at least 25 amino acids. In other embodiments, the purified polypeptide comprises fewer than 100 or 50 amino acids.

- In another embodiment, the purified polypeptide consists of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235. In another 5 embodiment, the purified polypeptide consists essentially of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

- Other embodiments include a polypeptide that contains one or more changes in amino acid sequence, e.g., a change in an amino acid residue that is not essential for activity, e.g., the ability of the polypeptide to bind to a MHC molecule or to be 10 recognized by an antibody described herein. Such polypeptides differ in amino acid sequence from SEQ ID NOs: 1-235, yet retain biological activity. In one embodiment, the polypeptide includes an amino acid sequence at least about 80%, 85%, 90%, 95%, 98% or more identical to any of SEQ ID NOs: 1-235. In another embodiment the polypeptide comprises an amino acid sequence selected from the group consisting of a 15 variant of any one of SEQ ID NOs: 1-235, wherein the variant has no more than two conservative amino acid substitutions. In another embodiment, the polypeptide comprises at least an immunogenic portion of a protein, wherein the protein comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

- The amino acid residues at particular positions in a polypeptide may include 20 analogs, derivatives and congeners of any specific amino acid referred to herein. For example, the present invention contemplates the use of amino acid analogs wherein a side chain is lengthened or shortened while still providing a carboxyl, amino or other reactive precursor functional group for cyclization, as well as amino acid analogs having variant side chains with appropriate functional groups. For instance, the subject polypeptide can 25 include an amino acid analog such as β -cyanoalanine, canavanine, djenkolic acid, norleucine, 3-phosphoserine, homoserine, dihydroxyphenylalanine, 5-hydroxytryptophan, 1-methylhistidine, or 3-methylhistidine. Other naturally occurring amino acid metabolites or precursors having side chains that are suitable herein will be recognized by those skilled in the art and are included in the scope of the present invention. Analogs of 30 polypeptides can be generated by mutagenesis, such as by discrete point mutation(s), or by truncation. For instance, mutation can give rise to analogs that retain substantially the

same, or merely a subset, of the biological activity of the polypeptide from which it was derived.

The polypeptides that can be utilized in the present invention also include analogs that are resistant to proteolytic cleavage such as those that, due to mutations, alter 5 ubiquitination or other enzymatic targeting associated with the protein.

Polypeptide analogs may also be chemically modified to create derivatives by forming covalent or aggregate conjugates with other chemical moieties, such as glycosyl groups, lipids, phosphate, acetyl groups and the like. Covalent derivatives of proteins can be prepared by linking the chemical moieties to functional groups on amino acid 10 sidechains of the protein or at the N-terminus or at the C-terminus of the polypeptide.

Modification of the structure of the subject polypeptides can be for such purposes as enhancing stability (e.g., *ex vivo* shelf life and resistance to proteolytic degradation *in vivo*), or post-translational modifications (e.g., to alter the phosphorylation pattern of the polypeptide). Such modified peptides, when designed to retain at least one activity of a 15 naturally-occurring form of the polypeptides disclosed herein, are considered to be their functional equivalents. Such modified peptides can be produced, for instance, by amino acid substitution, deletion, or addition.

For example, it is reasonable to expect that an isolated replacement of a leucine with an isoleucine or valine, an aspartate with a glutamate, a threonine with a serine, or a 20 similar replacement of an amino acid with a structurally related amino acid (i.e. isosteric and/or isoelectric mutations) will not have a major effect on the biological activity of the resulting molecule. Thus, altered nucleic acid sequences encoding polypeptides which are encompassed by the invention include deletions, insertions, or substitutions of different nucleotides resulting in a polynucleotide that encodes the same or a functionally 25 equivalent polypeptide. The encoded protein may also contain deletions, insertions, or substitutions of amino acid residues which produce a silent change and result in a functionally equivalent polypeptide. Deliberate amino acid substitutions may be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity, and/or the amphipathic nature of the residues as long as the biological activity of the 30 polypeptide is retained. Conservative replacements are those that take place within a family of amino acids that are related in their side chains. Genetically encoded amino

acids are can be divided into four families: (1) acidic=aspartate, glutamate; (2) basic=lysine, arginine, histidine; (3) nonpolar=alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan; and (4) uncharged polar=glycine, asparagine, glutamine, cysteine, serine, threonine, tyrosine (see, e.g., Biochemistry, 2nd 5 ed., Ed. by L. Stryer, W H Freeman and Co.: 1981). Whether a change in the amino acid sequence of a peptide results in a functional analog (e.g., functional in the sense that the resulting polypeptide mimics the wild-type form) can be readily determined by assessing the ability of the variant peptide to produce a response in cells in a fashion similar to the wild-type protein. Polypeptides in which more than one replacement has taken place can 10 readily be tested in the same manner.

As set forth above, alterations in primary sequence include genetic variations, both natural and induced. Also included are analogs that include residues other than naturally occurring L-amino acids, e.g., D-amino acids or non-naturally occurring or synthetic amino acids, e.g., β or δ amino acids. Alternatively, increased stability or 15 solubility may be conferred by cyclizing the peptide molecule.

A polypeptide of the invention preferably does not contain a peptide sequence described in Tables 1-10 of U.S. Patent No. 5,827,516.

The invention also features purified nucleic acids comprising nucleotides encoding polypeptides comprising amino acid sequences selected from the group 20 consisting of SEQ ID NOS: 1-235 or an amino acid sequence which is at least 95% identical to an amino acid sequence selected from the group consisting of SEQ ID NOS: 1-235. In another embodiment, the isolated nucleic acid comprises a nucleotide sequence that encodes a polypeptide comprising an amino acid sequence selected from the group 25 consisting of a variant of any one of SEQ ID NOS:1-235, wherein the variant has no more than two conservative amino acid substitutions.

In another embodiment, the isolated nucleic acid comprises a nucleotide sequence encoding a polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOS:1-235. In another embodiment, the isolated nucleic acid comprises a nucleotide sequence that encodes a polypeptide comprising at least 8 30 contiguous amino acids of an amino acid sequence selected from the group consisting of SEQ ID NOS:1-235. In another embodiment, the isolated nucleic acid comprises a

nucleotide sequence encoding a polypeptide consisting essentially of an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235.

In a further embodiment, the isolated nucleic acid comprises a nucleotide sequence encoding a polypeptide comprising no more than 30 contiguous amino acids of 5 a naturally occurring human protein, wherein the naturally occurring protein comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

A nucleic acid encoding a polypeptide described herein can be cloned into an expression vector, e.g., a vector in which the coding sequence is operably linked to expression control sequences. The need for, and identity of, expression control 10 sequences will vary according to the type of cell in which the DNA is to be expressed. Generally, expression control sequences can include any or all of the following: a transcriptional promoter, enhancer, suitable mRNA ribosomal binding sites, translation start site, and sequences that terminate transcription and translation, including polyadenylation and possibly translational control sequences. Suitable expression control 15 sequences can be selected by one of ordinary skill in the art. In one example, the vector comprises an expression control sequence that directs the expression of the polypeptides described herein. In another example, the vector comprises expression control sequences that direct expression of the nucleic acid molecule, as described herein. The nucleic acids encoding the polypeptides described herein may encode a methionine residue at the 20 amino terminus of the polypeptide to facilitate translation. Standard methods can be used by the skilled person to construct expression vectors. *See generally, Sambrook et al., 1989, Molecular Cloning: A Laboratory Manual (2nd Edition), Cold Spring Harbor Press, N.Y.*

Vectors useful in this invention include linear DNA with transcriptional control 25 elements, RNA, plasmid vectors, viral vectors, and bacterial vectors. A "plasmid" is an autonomous, self-replicating, extrachromosomal, circular DNA. Preferred viral vectors are those derived from retroviruses, adenovirus, adeno-associated virus, pox viruses, SV40 virus, alpha viruses or herpes viruses.

Isolated nucleic acids can be used for the *in vitro* production of polypeptides of 30 the invention. For example, a cell or cell line can be transfected, transformed, or infected with a nucleic acid described herein. After an incubation period that permits expression

of a polypeptide encoded by the nucleic acid, the polypeptide can be purified from the cell culture media, if secreted, or from a lysate of the cells expressing the polypeptide.

Fusion Proteins

5 The invention also provides fusion proteins. A "fusion protein" refers to a polypeptide containing a peptide sequence described herein, e.g., a peptide of any of SEQ ID NOs:1-235, and a heterologous amino acid sequence. A "heterologous amino acid sequence" refers to a sequence of contiguous amino acids that is not contained within the protein from which the peptide sequence is derived, e.g., a naturally occurring protein
10 that contains any of SEQ ID NOs:1-235. In other words, a fusion protein is not identical to a naturally occurring protein because it contains both a peptide sequence described herein as well as an amino acid sequence not contained within the naturally occurring protein from which the peptide sequence is derived. The fusion protein can contain a heterologous amino acid sequence fused to the N-terminus and/or C-terminus of the
15 peptide sequence.

The fusion protein can include a moiety that has a high affinity for a ligand. Such fusion proteins, e.g., GST-fusion proteins, can facilitate the purification of recombinant polypeptide. Fusion proteins can include all or a part of a serum protein, e.g., an IgG constant region, or human serum albumin.

20 The fusion protein can include a trafficking sequence. A "trafficking sequence" is an amino acid sequence that causes a polypeptide to which it is fused to be transported to a specific compartment of the cell. An example of a trafficking sequence is a signal sequence. In certain host cells (e.g., mammalian host cells), expression and/or secretion of a polypeptide can be increased through use of a heterologous signal sequence. For
25 example a signal sequence can be linked, with or without a linker, to a polypeptide described herein, e.g., a peptide of any of SEQ ID NOs:1-235.

Fusion proteins of the invention can be used as immunogens. For example, administration of a fusion protein, or a nucleic acid encoding a fusion protein, can be used to elicit an immune response in a host, e.g., a mammal such as a mouse, rat, or
30 human. Thus, the invention features an immunogenic composition comprising a polypeptide as described herein, the composition when injected into a mammal elicits an

immunogenic response directed against a polypeptide as described herein. The immunogenic response can be elicited by fragments of the polypeptide or nucleic acids encoding fragments of the polypeptide. Such fusion proteins may be useful in the development of antibodies, as described below.

5

Antibodies

The invention also includes an antibody, multispecific antibodies (e.g., bispecific antibodies), or a fragment thereof (e.g., an antigen-binding fragment thereof) that is specific for a peptide sequence described herein, e.g., a peptide of any of SEQ ID NOs:1-10 235. The term "antibody" as used herein refers to an immunoglobulin molecule or immunologically active portion thereof, i.e., an antigen-binding portion, including heterologous and chimeric antibodies.. The antibody can be a polyclonal or a monoclonal antibody. In other embodiments, the antibody can be recombinantly produced, e.g., produced by phage display or by combinatorial methods. "Antibody 15 fragments" comprise a portion of a full length antibody, generally the antigen binding or variable region thereof. Examples of antibody fragments include Fab, Fab', F(ab')₂, and Fv fragments and fragments produced by a Fab expression library; diabodies; linear antibodies; single-chain antibody molecules; and multispecific antibodies formed from antibody fragments. Neutralizing antibodies, (i.e., those which inhibit dimer formation) 20 are especially preferred for therapeutic use. Various techniques have been developed for the production of antibody fragments. Traditionally, these fragments were derived via proteolytic digestion of intact antibodies (see, e.g., Morimoto et al., Journal of Biochemical and Biophysical Methods 24: 107-117 (1992) and Brennan et al., Science 229: 81 (1985)). However, these fragments can now be produced directly by recombinant 25 host cells. For example, Fab'-SH fragments can be directly recovered from E. coli and chemically coupled to form F(ab')₂ fragments (Carter et al., Bio/Technology 10: 163-167 (1992)). According to another approach, F(ab')₂ fragments can be isolated directly from recombinant host cell culture. F(ab')₂ fragments can be produced by pepsin digestion of the antibody molecule and the Fab fragments which can be generated by reducing the 30 disulfide bridges of the F(ab')₂ fragments. Alternatively, Fab expression libraries may be constructed to allow rapid and easy identification of monoclonal Fab fragments with the

desired specificity (Huse, W. D. et al. (1989) Science 254:1275-1281). Other techniques for the production of antibody fragments will be apparent to the skilled practitioner.

- It may be desirable to generate multispecific (e.g. bispecific) humanized antibodies, as described herein, having binding specificities for at least two different epitopes. Exemplary bispecific antibodies may bind to two different epitopes of a protein. Alternatively, an arm may be combined with an arm which binds to a triggering molecule on a leukocyte such as a T-cell receptor molecule (e.g., CD2 or CD3), or Fc receptors for IgG (Fc. γ .R), such as Fc. γ .RI (CD64), Fc γ RII (CD32) and Fc. γ .RIII (CD16) so as to focus cellular defense mechanisms to the protein expressing cell.
- 10 Bispecific antibodies may also be used to localize cytotoxic agents to cells which express a protein. These antibodies possess a protein-binding arm and an arm which binds the cytotoxic agent (e.g., saporin, anti-interferon-.alpha., vinca alkaloid, ricin A chain, methotrexate or radioactive isotope hapten). Bispecific antibodies can be prepared as full length antibodies or antibody fragments (e.g., F(ab')₂ bispecific antibodies).
- 15 According to another approach for making bispecific antibodies, the interface between a pair of antibody molecules can be engineered to maximize the percentage of heterodimers which are recovered from recombinant cell culture. The preferred interface comprises at least a part of the C_H 3 domain of an antibody constant domain. In this method, one or more small amino acid side chains from the interface of the first antibody molecule are replaced with larger side chains (e.g., tyrosine or tryptophan).
- 20 Compensatory "cavities" of identical or similar size to the large side chain(s) are created on the interface of the second antibody molecule by replacing large amino acid side chains with smaller ones (e.g., alanine or threonine). This provides a mechanism for increasing the yield of the heterodimer over other unwanted end-products such as homodimers. See WO96/27011 published Sep. 6, 1996.
- 25 Bispecific antibodies include cross-linked or "heteroconjugate" antibodies. For example, one of the antibodies in the heteroconjugate can be coupled to avidin, the other to biotin. Heteroconjugate antibodies may be made using any convenient cross-linking methods. Suitable cross-linking agents are well known in the art, and are disclosed in U.S. Pat. No. 4,676,980, along with a number of cross-linking techniques.

Techniques for generating bispecific antibodies from antibody fragments have also been described in the literature. For example, bispecific antibodies can be prepared using chemical linkage. Brennan et al., *Science* 229: 81 (1985) describe a procedure wherein intact antibodies are proteolytically cleaved to generate F(ab')₂ fragments. These 5 fragments are reduced in the presence of the dithiol complexing agent sodium arsenite to stabilize vicinal dithiols and prevent intermolecular disulfide formation. The Fab' fragments generated are then converted to thionitrobenzoate (TNB) derivatives. One of the Fab'-TNB derivatives is then reconverted to the Fab'-thiol by reduction with mercaptoethylamine and is mixed with an equimolar amount of the other Fab'-TNB 10 derivative to form the bispecific antibody. The bispecific antibodies produced can be used as agents for the selective immobilization of enzymes.

Fab'-SH fragments can be recovered from *E. coli*, which can be chemically coupled to form bispecific antibodies. Shalaby et al., *J. Exp. Med.* 175: 217-225 (1992) describe the production of a fully humanized bispecific antibody F(ab')₂ molecule. Each 15 Fab' fragment was separately secreted from *E. coli* and subjected to directed chemical coupling *in vitro* to form the bispecific antibody. The bispecific antibody thus formed was able to bind to cells overexpressing the HER2 receptor and normal human T cells, as well as trigger the lytic activity of human cytotoxic lymphocytes against human breast tumor targets

20 Various techniques for making and isolating bispecific antibody fragments directly from recombinant cell culture have also been described. For example, bispecific antibodies have been produced using leucine zippers. Kostelny et al., *J. Immunol.* 148(5): 1547-1553 (1992). The leucine zipper peptides from the Fos and Jun proteins were linked to the Fab' portions of two different antibodies by gene fusion. The antibody homodimers 25 were reduced at the hinge region to form monomers and then re-oxidized to form the antibody heterodimers. This method can also be utilized for the production of antibody homodimers. The "diabody" technology described by Hollinger et al., *Proc. Natl. Acad. Sci. USA* 90: 6444-6448 (1993) has provided an alternative mechanism for making bispecific antibody fragments. The fragments comprise a heavy-chain variable domain 30 (V_H) connected to a light-chain variable domain (V_L) by a linker which is too short to allow pairing between the two domains on the same chain. Accordingly, the V_H and V_L

domains of one fragment are forced to pair with the complementary V_L and V_H domains of another fragment, thereby forming two antigen-binding sites. Another strategy for making bispecific antibody fragments by the use of single-chain Fv (sFv) dimers has also been reported. See Gruber et al., J. Immunol. 152: 5368 (1994). Alternatively, the
5 bispecific antibody may be a "linear antibody" produced as described in Zapata et al.
Protein Eng. 8(10): 1057-1062 (1995).

The term "monoclonal antibody" as used herein refers to an antibody obtained from a population of substantially homogeneous antibodies, i.e., the individual antibodies comprising the population are identical except for possible naturally occurring mutations
10 that may be present in minor amounts. Monoclonal antibodies are highly specific, being directed against a single antigenic site. Furthermore, in contrast to conventional (polyclonal) antibody preparations which typically include different antibodies directed against different determinants (epitopes), each monoclonal antibody is directed against a single determinant on the antigen. The modifier "monoclonal" indicates the character of
15 the antibody as being obtained from a substantially homogeneous population of antibodies, and is not to be construed as requiring production of the antibody by any particular method. For example, the monoclonal antibodies to be used in accordance with the present invention may be made by the hybridoma method first described by Kohler et al., Nature 256: 495 (1975), or may be made by recombinant DNA methods (see, e.g.,
20 U.S. Pat. No. 4,816,567). The "monoclonal antibodies" may also be isolated from phage antibody libraries using the techniques described in Clackson et al., Nature 352: 624-628 (1991) and Marks et al., J. Mol. Biol. 222: 581-597 (1991), for example.

The monoclonal antibodies herein specifically include "chimeric" antibodies (immunoglobulins) in which a portion of the heavy and/or light chain is identical with or
25 homologous to corresponding sequences in antibodies derived from a particular species or belonging to a particular antibody class or subclass, while the remainder of the chain(s) is identical with or homologous to corresponding sequences in antibodies derived from another species or belonging to another antibody class or subclass, as well as fragments of such antibodies, so long as they exhibit the desired biological activity
30 (U.S. Pat. No. 4,816,567; and Morrison et al., Proc. Natl. Acad Sci. USA 81: 6851-6855 (1984)).

Polyclonal antibodies are preferably raised in animals by multiple subcutaneous (sc) or intraperitoneal (ip) injections of the relevant antigen and an adjuvant. It may be useful to conjugate the relevant antigen to a protein that is immunogenic in the species to be immunized, e.g., keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, or 5 soybean trypsin inhibitor using a bifunctional or derivatizing agent, for example, maleimidobenzoyl sulfosuccinimide ester (conjugation through cysteine residues), N-hydroxysuccinimide (through lysine residues), glutaraldehyde, succinic anhydride, SOCl_2 , or $\text{R}^1\text{N}=\text{C}=\text{NR}$, where R and R^1 are different alkyl groups.

A polypeptide described herein, e.g., a peptide of any of SEQ ID NOs:1-235, can 10 be used as an immunogen or can be used to identify antibodies made with other immunogens, e.g., cells, membrane preparations, and the like. Polypeptides can be expressed on the cell surface enabling the binding of an antibody, as described herein, that is specific to the polypeptide. Alternately, an antibody described herein may bind to a polypeptide described herein, where the polypeptide is a target of a second antibody 15 located on the cell surface.

An antibody (e.g., a monoclonal antibody) can be used to isolate a polypeptide described herein by standard techniques, such as affinity chromatography or immunoprecipitation. Moreover, an antibody can be used to detect the polypeptide (e.g., 20 in a cellular lysate or cell supernatant) in order to evaluate the abundance and pattern of expression of the protein.

Furthermore, an antibody can be used to target a protein *in vivo* for a variety of purposes including disease screening, diagnosis, and treatment. For example, an antibody can be modified to include a toxin and/or a detectable label, as described herein. Antibodies coupled to a toxic agent can be particularly useful to target and destroy 25 diseased or infected cells.

An antibody can be coupled to a toxin, e.g., a polypeptide toxin, e.g., ricin or diphtheria toxin or active fragment thereof, or a radioactive nucleus, or imaging agent, e.g. a radioactive, enzymatic, or other, e.g., imaging agent, e.g., a NMR contrast agent. Toxins can be optionally in an inactive state and be subject to activation following their 30 administration to a subject (e.g., activation via radio energy, irradiation with x-rays, or other penetrating rays). Labels which produce detectable radioactive emissions or

fluorescence are preferred. Examples of detectable substances that can be coupled to an antibody include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, and radioactive materials.

- "Humanized" forms of non-human (e.g., murine) antibodies are chimeric
- 5 antibodies which contain minimal sequence derived from non-human immunoglobulin. For the most part, humanized antibodies are human immunoglobulins (recipient antibody) in which hypervariable region residues of the recipient are replaced by hypervariable region residues from a non-human species (donor antibody) such as mouse, rat, rabbit or nonhuman primate having the desired specificity, affinity, and capacity.
- 10 The resulting antibody is one in which amino acids have been replaced in the non-antigen binding regions in order to more closely resemble a human antibody, while still retaining the original binding ability. In some instances, Fv framework region (FR) residues of the human immunoglobulin are replaced by corresponding non-human residues. Furthermore, humanized antibodies may comprise residues which are not found in the
- 15 recipient antibody or in the donor antibody. These modifications are made to further refine antibody performance. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the hypervariable loops correspond to those of a non-human immunoglobulin and all or substantially all of the FR regions are those of a human immunoglobulin sequence. The humanized antibody optionally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin. For further details, see Jones et al., Nature 321: 522-525 (1986); Reichmann et al., Nature 332: 323-329 (1988); and Presta, Curr. Op. Struct. Biol. 2: 593-596 (1992).
- 20 "Single-chain Fv" or "sFv" antibody fragments comprise the V_H and V_L domains of antibody, wherein these domains are present in a single polypeptide chain. Generally, the Fv polypeptide further comprises a polypeptide linker between the V_H and V_L domains which enables the sFv to form the desired structure for antigen binding. For a review of sFv see Pluckthun in The Pharmacology of Monoclonal Antibodies, vol. 113,
- 25 Rosenburg and Moore eds. Springer-Verlag, New York, pp. 269-315 (1994).

Humanized antibodies can be produced, for example by transgenic non-human animals. Such animals are capable of producing heterologous antibodies of multiple isotypes. Heterologous antibodies are encoded by immunoglobulin heavy chain genes not normally found in the genome of that species of non-human animal. Transgenic non-
5 human animals (e.g., mammals) can be of a variety of species including murine (rodents (e.g., mice, rats), avian (chicken, turkey, fowl), bovine (beef, cow, cattle), ovine (lamb, sheep, goats), porcine (pig, swine), and piscine (fish). Transgenic non-human animals can be produced by introducing transgenes into the germline of the non-human animal.
A "transgene" means a nucleic acid sequence (encoding, e.g., a human Fc receptor),
10 which is partly or entirely heterologous, i.e., foreign, to the transgenic animal or cell into which it is introduced, or, is homologous to an endogenous gene of the transgenic animal or cell into which it is introduced, but which is designed to be inserted, or is inserted, into the animal's genome in such a way as to alter the genome of the cell into which it is inserted (e.g., it is inserted at a location which differs from that of the natural gene or its
15 insertion results in a knockout). A transgene can include one or more transcriptional regulatory sequences and any other nucleic acid, such as introns, that may be necessary for optimal expression of a selected nucleic acid. Methods of producing transgenic animals and humanized antibodies are for example described in U.S. patents 5,569,825, 5,770,429, and 6,11,166.

20 Humanized antibodies can be bound to labels or be in the form of a conjugate bound to a cytotoxic agent. The word "label" when used herein refers to a detectable compound or composition which is conjugated directly or indirectly to the antibody. The label may itself be detectable by itself (e.g., radioisotope labels or fluorescent labels) or, in the case of an enzymatic label, may catalyze chemical alteration of a substrate
25 compound or composition which is detectable. The term "cytotoxic agent" as used herein refers to a substance that inhibits or prevents the function of cells and/or causes destruction of cells. The term is intended to include radioactive isotopes (e.g., I.¹³¹, I.¹²⁵, Y⁹⁰ and Re¹⁸⁶), chemotherapeutic agents, and toxins such as enzymatically active toxins of bacterial, fungal, plant or animal origin, or fragments thereof. A "chemotherapeutic
30 agent" is a chemical compound useful in the treatment of cancer. Examples of chemotherapeutic agents include Adriamycin, Doxorubicin, 5-Fluorouracil, Cytosine

arabinoside ("Ara-C"), Cyclophosphamide, Thiotepa, Taxotere (docetaxel), Busulfan, Cytoxin, Taxol, Methotrexate, Cisplatin, Melphalan, Vinblastine, Bleomycin, Etoposide, Ifosfamide, Mitomycin C, Mitoxantrone, Vincreistine, Vinorelbine, Carboplatin, Teniposide, Daunomycin, Carminomycin, Aminopterin, Dactinomycin, Mitomycins, 5 Esperamicins (see U.S. Pat. No. 4,675,187), Melphalan and other related nitrogen mustards.

Covalent modifications of the humanized antibody are also included within the scope of this invention. They may be made by chemical synthesis or by enzymatic or chemical cleavage of the antibody, if applicable. Other types of covalent modifications of 10 the antibody are introduced into the molecule by reacting targeted amino acid residues of the antibody with an organic derivatizing agent that is capable of reacting with selected side chains or the N- or C-terminal residues.

Cysteinyl residues most commonly are reacted with .alpha.-haloacetates (and corresponding amines), such as chloroacetic acid or chloroacetamide, to give 15 carboxymethyl or carboxyamidomethyl derivatives. Cysteinyl residues also are derivatized by reaction with bromotrifluoroacetone, .alpha.-bromo-.beta.-(5-imidozoyl)propionic acid, chloroacetyl phosphate, N-alkylmaleimides, 3-nitro-2-pyridyl disulfide, methyl 2-pyridyl disulfide, p-chloromercuribenzoate, 2-chloromercuri-4-nitrophenol, or chloro-7-nitrobenzo-2-oxa-1,3-diazole.

20 Histidyl residues are derivatized by reaction with diethylpyrocarbonate at pH 5.5-7.0 because this agent is relatively specific for the histidyl side chain. Para-bromophenacyl bromide also is useful; the reaction is preferably performed in 0.1 M sodium cacodylate at pH 6.0.

Lysinyl and amino-terminal residues are reacted with succinic or other carboxylic 25 acid anhydrides. Derivatization with these agents has the effect of reversing the charge of the lysinyl residues. Other suitable reagents for derivatizing .alpha.-amino-containing residues include imidoesters such as methyl picolinimidate, pyridoxal phosphate, pyridoxal, chloroborohydride, trinitrobenzenesulfonic acid, O-methylisourea, 2,4-pentanedione, and transaminase-catalyzed reaction with glyoxylate.

30 Arginyl residues are modified by reaction with one or several conventional reagents, among them phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, and

ninhydrin. Derivatization of arginine residues requires that the reaction be performed in alkaline conditions because of the high pKa of the guanidine functional group.

Furthermore, these reagents may react with the groups of lysine as well as the arginine epsilon-amino group.

5 The specific modification of tyrosyl residues may be made, with particular interest in introducing spectral labels into tyrosyl residues by reaction with aromatic diazonium compounds or tetranitromethane. Most commonly, N-acetylimidazole and tetranitromethane are used to form O-acetyl tyrosyl species and 3-nitro derivatives, respectively. Tyrosyl residues are iodinated using ^{125}I or ^{131}I to prepare labeled proteins
10 for use in radioimmunoassay.

Carboxyl side groups (aspartyl or glutamyl) are selectively modified by reaction with carbodiimides ($\text{R}-\text{N}=\text{C}=\text{N}-\text{R}'$), where R and R' are different alkyl groups, such as 1-cyclohexyl-3-(2-morpholinyl-4-ethyl) carbodiimide or 1-ethyl-3-(4-azonia-4,4-dimethylpentyl) carbodiimide. Furthermore, aspartyl and glutamyl residues are converted
15 to asparaginyl and glutaminyl residues by reaction with ammonium ions.

Glutaminyl and asparaginyl residues are frequently deamidated to the corresponding glutamyl and aspartyl residues, respectively. These residues are deamidated under neutral or basic conditions. The deamidated form of these residues falls within the scope of this invention.

20 Other modifications include hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of seryl or threonyl residues, methylation of the .alpha.-amino groups of lysine, arginine, and histidine side chains (T. E. Creighton, Proteins: Structure and Molecular Properties, W.H. Freeman & Co., San Francisco, pp. 79-86 (1983)), acetylation of the N-terminal amine, and amidation of any C-terminal carboxyl group.

25 Another type of covalent modification involves chemically or enzymatically coupling glycosides to the antibody. These procedures are advantageous in that they do not require production of the antibody in a host cell that has glycosylation capabilities for N- or O-linked glycosylation. Depending on the coupling mode used, the sugar(s) may be attached to (a) arginine and histidine, (b) free carboxyl groups, (c) free sulphydryl groups
30 such as those of cysteine, (d) free hydroxyl groups such as those of serine, threonine, or hydroxyproline, (e) aromatic residues such as those of phenylalanine, tyrosine, or

tryptophan, or (f) the amide group of glutamine. These methods are described in WO87/05330 published 11 Sep. 1987, and in Aplin and Wriston, CRC Crit. Rev. Biochem., pp. 259-306 (1981).

Removal of any carbohydrate moieties present on the antibody may be
5 accomplished chemically or enzymatically. Chemical deglycosylation requires exposure of the antibody to the compound trifluoromethanesulfonic acid, or an equivalent compound. This treatment results in the cleavage of most or all sugars except the linking sugar (N-acetylglucosamine or N-acetylgalactosamine), while leaving the antibody intact. Chemical deglycosylation is described by Hakimuddin, et al. Arch. Biochem. Biophys.
10 259: 52 (1987) and by Edge et al. Anal. Biochem., 118: 131 (1981). Enzymatic cleavage of carbohydrate moieties on antibodies can be achieved by the use of a variety of endo- and exo-glycosidases as described by Thotakura et al. Meth. Enzymol. 138: 350 (1987).

Another type of covalent modification of the antibody comprises linking the antibody to one of a variety of nonproteinaceous polymers, e.g., polyethylene glycol,
15 polypropylene glycol, or polyoxyalkylenes, in the manner set forth in U.S. Pat. Nos. 4,640,835; 4,496,689; 4,301,144; 4,670,417; 4,791,192 or 4,179,337.

Humanized antibodies can also be immobilized to a solid phase. By "solid phase" is meant a non-aqueous matrix to which the antibody of the present invention can adhere. Examples of solid phases encompassed herein include those formed partially or entirely
20 of glass (e.g. controlled pore glass), polysaccharides (e.g., agarose), polyacrylamides, polystyrene, polyvinyl alcohol and silicones. In certain embodiments, depending on the context, the solid phase can comprise the well of an assay plate; in others it is a purification column (e.g. an affinity chromatography column). This term also includes a discontinuous solid phase of discrete particles, such as those described in U.S. Pat. No.
25 4,275,149.

Diagnostic and therapeutic uses for the antibody are contemplated. In one diagnostic application, the invention provides a method for determining the presence of a protein comprising exposing a sample suspected of containing the protein to the antibody and determining binding of the antibody to the sample. For this use, the invention
30 provides a kit comprising the antibody and instructions for using the antibody to detect the protein.

Also included in the invention is an isolated nucleic acid, as described herein, encoding a humanized antibody, described herein, as well as a vector comprising the nucleic acid and a cell comprising the vector.

- Antibodies can be used to modulate the activity of a polypeptide of the invention,
- 5 as described herein. The invention includes a method for modulating the activity of the polypeptide of the invention, the method comprising contacting the polypeptide with a compound that binds to the polypeptide in a concentration sufficient to modulate the activity of the polypeptide. The compound that binds to the polypeptide can be an antibody as described herein.
- 10 The invention also features a method of making an antibody, the method comprising (a) providing a polypeptide described herein to a mammal in an amount effective to induce the production of an antibody that binds to the polypeptide; (b) isolating from the mammal a cell that produces an antibody that selectively binds to a polypeptide as described herein; (c) immortalizing the cell isolated in step (b); and (d) isolating antibodies from the immortalized
- 15 cell.

The invention also includes a method of selecting an antibody, the method comprising: (a) contacting a polypeptide as described herein with an in vitro library of antibodies; (b) binding an antibody to the polypeptide; and (c) selecting the antibody that binds to the polypeptide.

- 20 The invention also includes a nucleic acid that encodes an antibody described herein. Also included are vectors that include the nucleic acid and cells transformed with the nucleic acid, particularly cells which are useful for producing an antibody, e.g., mammalian cells, e.g. CHO or lymphatic cells.

The invention also includes cell lines, e.g., hybridomas, which make an antibody

25 described herein, and method of using said cells to make an antibody.

- Also included in the invention are anti-peptide antibodies. An anti-peptide antibody is an antibody that binds to the amino acid sequence of a peptide described herein, e.g., a peptide of any of SEQ ID NOS:1-235. In one example, the antibody is capable of recognizing the peptide when the peptide is bound to an MHC class I or
- 30 class II molecule. The antibody can recognize either the peptide sequence or a

combination of the peptide sequence and an MHC molecule. See, e.g., Apostolopoulos *et al.*, 1998, *J. Immunol.* 161:767 for a description of anti-peptide antibodies.

The anti-peptide antibodies can be used to detect the expression of a protein within a cell (e.g., detection of a processed peptide on the cell surface by an anti-peptide antibody indicates that the protein, e.g., intracellular protein, is expressed within the cell). Such an anti-peptide antibody can be particularly useful for determining the protein composition of a cell when the cell is subjected to varying conditions or stimuli. Additionally, an anti-peptide antibody can be useful for detecting the presence of a disease-associated antigen within a cell. For example, a cell can be diagnosed as containing a cancer-related protein by detecting a peptide described herein presented by an MHC molecule on the surface of the cell. Antibodies raised against peptides can also be used therapeutically to treat human maladies. For example, such an antibody can be modified to contain a reagent, e.g., a toxin, that damages or destroys diseased or infected cells to which it binds.

15

Gene Discovery

The human genome has been reported to contain approximately 30,000-40,000 genes, a number significantly lower than previous estimates of 100,000 or more genes.

Venter *et al.*, *Science* 2001 291:1304; International Human Genome Sequencing

20 Consortium *Nature* 2001 409:860. One possible explanation for this discrepancy is that computer algorithms used to analyze raw nucleotide sequence and identify genes may not have detected a subset of the genes in the human genome. Because the peptides described herein correspond to portions of actual proteins actually produced by a cell, the compositions and methods of the invention allow for the identification of as yet 25 unidentified genes. For example, those peptides that do not match to any known genes may represent the protein product of a novel gene.

A peptide sequence described herein can be compared to a predicted translation of human genomic sequence (a predicted translation of each strand of genomic DNA, in three reading frames). If this analysis identifies a matching sequence, then a careful 30 analysis of the reading frame encoding the peptide should allow for identification of the remainder of the gene encoding the peptide, including but not limited to coding

sequences, 5' and 3' untranslated regions, alternatively spliced exons, introns, promoters, enhancers, and silencer or repressor elements.

In addition to sequence analysis, a gene and/or a cDNA encoding a protein containing a peptide described herein can be isolated by methods well known to those of skill in the art. Isolation of a gene or a cDNA is especially relevant for peptides that lack a genomic match, but can also be useful to verify the nucleotide sequence that encodes any peptide. The skilled artisan will appreciate that a number of methods are known in the art to identify and isolate genes or cDNAs using amino acid information, and will know how to identify and practice such methods. See, for example, Sambrook *et al.*, 1989 Molecular Cloning: A Laboratory Manual 2nd ed. Cold Spring Harbor Laboratory Press; Ausubel *et al.*, Current Protocols in Molecular Biology, Greene Publishing Associates and Wiley Interscience, N.Y. (current edition). Such methods include the preparation of degenerate probes or primers based upon the peptide amino acid sequence and using such primers for identification and/or amplification of genes and or cDNAs in appropriate libraries or other sources of genomic materials. The chromosomal location of the gene encoding the protein from which a peptide is derived may be determined, for example, by hybridizing appropriately labeled nucleic acids to chromosomes *in situ*.

Detection of Protein Expression

The compositions and methods described herein can be used to determine the protein composition of a cell. The detection of mRNA within a cell, for example by Northern analysis or RT-PCR, does not indicate whether the mRNA is translated, much less how much of the corresponding protein is produced in the cell. Detection of a peptide described herein indicates that the protein from which it is derived has been produced by the cell. Thus, the invention includes a method of determining the protein composition of a cell (or tissue sample) by detecting the presence of a peptide described herein to thereby determine that the cell (or tissue sample) expresses the protein from which the peptide is derived. The method can be used to determine the presence of a peptide and/or the protein from which it is derived, and optionally the quantity of a peptide and/or protein produced by a cell.

In addition to the "translational verification" described above, the peptides can be used to determine the reading frame that is being used by a gene. For example, the detection of an mRNA or a portion of an mRNA does not automatically indicate the amino acid sequence of the corresponding protein. The peptides described herein can 5 thus be used to discover reading frames of genes that are being expressed.

Protein Classifications

The peptides described herein belong to a wide variety of functional biological classes. Many of the classes to which particular peptides belong are described in the 10 Table presented in the Examples. Members of many of these classes of proteins have been well-characterized as participating in important biological pathways and/or have been implicated in a variety of disease conditions. Several of these classes are described in more detail below.

15 Kinases

As described in Examples 1 and 2 (and the accompanying table), many of the peptides described herein are derived from proteins that appear to be kinases. Kinases catalyze the transfer of high energy phosphate groups from a phosphate donor to a phosphate acceptor. Nucleotides usually serve as the phosphate donor in these reactions, 20 with most kinases utilizing adenosine triphosphate (ATP). Reversible protein phosphorylation is a primary method for regulating protein activity in eukaryotic cells. In general, proteins are activated by phosphorylation in response to extracellular signals such as hormones, neurotransmitters, and growth and differentiation factors. The activated proteins initiate the cell's intracellular response by way of intracellular 25 signaling pathways and second messenger molecules, such as cyclic nucleotides, calcium-calmodulin, inositol, and various mitogens, that regulate protein phosphorylation.

Kinases are involved in many aspects of a cell's function, from basic metabolic processes such as glycolysis, to cell-cycle regulation, differentiation, and communication 30 with the extracellular environment through signal transduction cascades. Kinase targets include proteins, inositol, lipids, and nucleotides. Inappropriate phosphorylation of

proteins in cells has been linked to changes in cell cycle progression and cell differentiation. Changes in cell cycle progression have been linked to induction of apoptosis or cancer. Changes in cell differentiation have been linked to diseases and disorders of the reproductive system, immune system, and skeletal muscle.

- 5 Table 1 lists several of the peptides described herein that appear, based upon structural homology, to belong to the kinase superfamily. These peptides (and their corresponding source proteins) can therefore be used to treat disorders associated with inappropriate phosphorylation of kinase targets, e.g., disorders associated with changes in cell cycle progression and/or cell differentiation, or to screen for agonists and antagonists
- 10 useful for the same purpose. In addition, nucleic acids encoding the proteins as well as compounds (e.g., antibodies) that recognize the proteins can be used in a wide variety of applications described herein, including therapeutics, diagnostics, and drug screening.

The invention therefore includes the following peptides as kinases: SEQ ID NO:7, SEQ ID NO:12, SEQ ID NO:32, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:45, SEQ
15 ID NO:85, SEQ ID NO:90, SEQ ID NO:95, SEQ ID NO:118, SEQ ID NO:140, SEQ ID
NO:181, and SEQ ID NO:185.

Phosphatases

As described in Examples 1 and 2 (and the accompanying table), many of the
20 peptides described herein are derived from proteins that appear to be phosphatases. Phosphatases are characterized as tyrosine-specific or serine/threonine-specific based on their preferred phospho-amino acid substrate. Some phosphatases exhibit dual specificity for both phospho-tyrosine and phospho-serine/threonine residues.

Serine/threonine phosphatases play important roles in glycogen metabolism,
25 muscle contraction, protein synthesis, oocyte maturation, and hepatic metabolism. (Cohen, P. (1989) Annu. Rev. Biochem. 58:453-508). Tyrosine phosphatases play important roles in lymphocyte activation and cell adhesion. In addition, the genes encoding several tyrosine phosphatases have been mapped to chromosomal regions that are translocated or rearranged in various neoplastic conditions, including lymphoma,
30 leukemia, small cell lung carcinoma, adenocarcinoma, and neuroblastoma (Charbonneau, H. and Tonks, N. K. (1992) Annu. Rev. Cell Biol. 8:463-493). Because cellular

transformation is often accompanied by increased phosphorylation activity, the regulation of phosphorylation activity by phosphatases may therefore be an important strategy for controlling some types of cancer.

Table 1 lists several of the peptides described herein that appear, based upon structural homology, to belong to the phosphatase superfamily. These peptides (and their corresponding source proteins) can therefore be used to treat disorders associated with inappropriate phosphorylation and/or phosphatase activity, or to screen for agonists and antagonists useful for the same purpose. These protein phosphatases and the nucleic acids encoding them allow for the manufacture of new compositions that are useful in the diagnosis, prevention, and treatment of disorders such as immune system disorders, cell proliferative and differentiative disorders (including cancer), and neurological disorders.

The invention therefore includes the following peptides as phosphatases: SEQ ID NO:18, SEQ ID NO:24, SEQ ID NO:76, SEQ ID NO:103, SEQ ID NO:125, SEQ ID NO:199, SEQ ID NO:224, and SEQ ID NO:231

15

Proteases and Protease Inhibitors

As described in Examples 1 and 2 (and the accompanying table), many of the peptides described herein are derived from proteins that appear to be proteases. Proteases cleave proteins and peptides at the peptide bond that forms the backbone of the protein or peptide chain. Proteolytic processing is an essential component of cell growth, differentiation, remodeling, and homeostasis. The cleavage of peptide bonds within cells is necessary for the maturation of precursor proteins to their active forms, the removal of signal sequences from targeted proteins, the degradation of incorrectly folded proteins, and the controlled turnover of peptides within the cell.

25 Proteases participate in apoptosis (and disorders associated with inappropriate levels of apoptosis) as well as tissue remodeling during embryonic development, wound healing, and normal growth. Proteases are involved in the etiology or progression of disease states such as inflammation, angiogenesis, tumor dispersion and metastasis, cardiovascular disease, neurological disease, and bacterial, parasitic, and viral infections.

30 For example, caspases and components of caspase signaling pathways regulate apoptosis and/or inflammation in an individual.

Protease inhibitors and other regulators of protease activity control the activity and effects of proteases. Protease inhibitors have been shown to control pathogenesis in animal models of proteolytic disorders and in the treatment of HIV (Murphy, G. (1991) Agents Actions Suppl. 35:69-76).

- 5 Table 1 lists several of the peptides described herein that appear, based upon structural homology, to be proteases. These peptides (and their corresponding source proteins) can therefore be used to treat disorders associated with inappropriate protease expression or activity. Examples of such disorder include immunological disorders (including autoimmune or inflammatory disorders), angiogenesis, tumor dispersion and
10 metastasis, cardiovascular disease, neurological disease, and pathogenic infections, or to screen for agonists and antagonists useful for the same purpose. In addition, nucleic acids encoding the proteins as well as compounds (e.g., antibodies) that recognize the proteins can be used in a wide variety of applications described herein, including therapeutics, diagnostics, and drug screening.
- 15 The invention therefore includes the following peptides as proteases: SEQ ID NO:75, SEQ ID NO:93, SEQ ID NO:163, SEQ ID NO:169, and SEQ ID NO:200.

Transporters

- As described in Examples 1 and 2 (and the accompanying table), many of the
20 peptides described herein are derived from proteins that appear to be transporters. Transporter proteins are used to facilitate the translocation of certain molecules either into or out of the cell. Often, such transporters work by "pumping" ions across the cell membrane and co-transporting specific molecules (e.g., amino acids, amino acid derivatives and precursors, dicarboxylates, or inorganic molecules) across the membrane.
25 Such mechanisms play important roles in maintaining cellular and metabolic homeostasis, neuron function, signaling, and drug resistance. As such, transporter proteins constitute compelling targets for the development of novel therapeutic agents.

- The electrical potential of a cell is generated and maintained by controlling the movement of ions across the plasma membrane. The movement of ions requires ion
30 channels, which form ion selective pores within the membrane. There are two basic types of ion channels, ion transporters and gated ion channels. Ion transporters utilize the

energy obtained from ATP hydrolysis to actively transport an ion against the ion's concentration gradient. Gated ion channels allow passive flow of an ion down the ion's electrochemical gradient under restricted conditions. Together, these types of ion channels generate, maintain, and utilize an electrochemical gradient that is used in 1) 5 electrical impulse conduction down the axon of a nerve cell, 2) transport of molecules into cells against concentration gradients, 3) initiation of muscle contraction, and 4) endocrine cell secretion.

The etiology of numerous human diseases and disorders can be attributed to defects in the transport of molecules across membranes. Defects in the trafficking of 10 membrane-bound transporters and ion channels are associated with several disorders, e.g., cystic fibrosis, glucose-galactose malabsorption syndrome, hypercholesterolemia, von Gierke disease, and certain forms of diabetes mellitus. Single-gene defect diseases resulting in an inability to transport small molecules across membranes include, e.g., cystinuria, iminoglycinuria, Hartup disease, and Fanconi disease (vant Hoff, W.G. (1996) 15 Exp. Nephrol. 4:253-262; Talente, G.M. et al. (1994) Ann. Intern. Med. 120:218-226; and Chillon, M. et al. (1995) New Engl. J. Med. 332:1475-1480).

Human diseases caused by mutations in ion channel genes include disorders of skeletal muscle, cardiac muscle, and the central nervous system. Mutations in the pore forming subunits of sodium and chloride channels cause myotonia, a muscle disorder in 20 which relaxation after voluntary contraction is delayed. Sodium channel myotonias have been treated with channel blockers. Mutations in muscle sodium and calcium channels cause forms of periodic paralysis, while mutations in the sarcoplasmic calcium release channel and muscle sodium channel cause malignant hyperthermia. Cardiac arrhythmia disorders such as the long QT syndromes and idiopathic ventricular fibrillation are caused 25 by mutations in potassium and sodium channels (Cooper, E.C. and L.Y. Jan (1998) Proc. Natl. Acad. Sci. USA 96:4759-4766). All four known human idiopathic epilepsy genes code for ion channel proteins (Berkovic, S.F. and I.E. Scheffer (1999) Curr. Opin. Neurology 12:177 182). Other neurological disorders such as ataxias and hereditary deafness can also result from mutations in ion channel genes (Jen, J. (1999) Curr. Opin. 30 Neurobiol. 9:274-280).

- Ion channels have been the target for many drug therapies. In particular, neurotransmitter-gated channels have been targeted in therapies for treatment of insomnia, anxiety, depression, and schizophrenia. Voltage-gated channels have been targeted in therapies for arrhythmia, ischemia, stroke, head trauma, and
- 5 neurodegenerative disease (Taylor, C.P. and L. S. Narasimhan (1997) *Adv. Pharmacol.* 39:47-98).

Various classes of ion channels also play an important role in the perception of pain, and thus are potential targets for new analgesics. These include the vanilloid-gated ion channels, which are activated by the vanilloid capsaicin, as well as by noxious heat.

10 Local anesthetics such as lidocaine and mexiletine which blockade voltage-gated ion channels have been useful in the treatment of neuropathic pain.

Ion channels in the immune system have been suggested as targets for immunomodulation. T-cell activation depends upon calcium signaling, and a diverse set of T-cell specific ion channels has been characterized that affect this signaling process.

15 Channel blocking agents can inhibit secretion of lymphokines, cell proliferation, and killing of target cells.

Table 1 lists several of the peptides described herein that appear, based upon structural homology, to be transporters. These peptides (and their corresponding source proteins) can therefore be used to treat disorders associated with inappropriate transporter expression or activity. Examples of such disorders include neurological, muscle, and immunological disorders, or to screen for agonists and antagonists useful for the same purpose. In addition, nucleic acids encoding the proteins as well as compounds (e.g., antibodies) that recognize the proteins can be used in a wide variety of applications described herein, including therapeutics, diagnostics, and drug screening.

20 The invention therefore includes the following peptides as transporters: SEQ ID NO:1, SEQ ID NO:25, SEQ ID NO:48, SEQ ID NO:51, SEQ ID NO:52, SEQ ID NO:56, SEQ ID NO:58, SEQ ID NO:59, SEQ ID NO:61, SEQ ID NO:62, SEQ ID NO:63, SEQ ID NO:64, SEQ ID NO:77, SEQ ID NO:78, SEQ ID NO:79, SEQ ID NO:80, SEQ ID NO:81, SEQ ID NO:82, SEQ ID NO:83, SEQ ID NO:84, SEQ ID NO:90, SEQ ID NO:94, SEQ ID NO:100, SEQ ID NO:116, SEQ ID NO:128, SEQ ID NO:130, SEQ ID NO:131, SEQ ID NO:132, SEQ ID NO:133, SEQ ID NO:141, SEQ ID NO:170, SEQ ID

NO:178, SEQ ID NO:187, SEQ ID NO:189, SEQ ID NO:203, SEQ ID NO:207, SEQ ID NO:219, and SEQ ID NO:234.

Cytoskeletal Proteins

5 As described in Examples 1 and 2 (and the accompanying table), many of the peptides described herein are derived from proteins that appear to be cytoskeletal proteins. The physical-biochemical processes of cell motility, organelle movement, chromosome movement, cytokinesis, and generation of cell shape are all dependent on a complex of protein fibers found in the cytoplasm. This protein complex is termed the
10 cytoskeleton. The cytoskeleton of eukaryotic cells has three major filamentous systems. These systems are the actin filaments, intermediate filaments, and microtubules. Each of these filamentous systems is assembled from different proteins, including actin, myosin, tubulins, and intermediate filament proteins. Different cell types and tissues express specific isoforms of the proteins which comprise these filaments. In some cases distinct
15 isoforms and mRNA splice variants are associated with cell-type specific functions (Lees-Miller, J.P. and Helfman, D.M. (1991) BioEssays 13:429-437).

Cell motility is governed by the interaction between cytoskeletal and other cellular proteins. Cytoskeletal proteins that are involved in the generation of motive force within the cell are termed contractile proteins. Cytoskeletal proteins are involved in
20 the regulation of muscle contraction. Vertebrate smooth muscle contraction is dependent upon levels of cAMP and intracellular calcium ions.

Cytoskeletal proteins are implicated in several diseases. Pathologies such as muscular dystrophy, nephrotic syndrome, and dilated cardiomyopathy have been associated with differential expression of alpha-actinin-3 (Vainzof, M. et al. (1997)
25 Neuropediatrics 28:223-228; Smoyer, W.E. and Mundel, P. (1998) J. Mol. Med. 76:172-183; and Sussman, M.A. et al. (1998) J. Clin. Invest. 101:51-61). Alpha actinin and several microtubule associated proteins (MAPs) are present in Hirano bodies, which are observed more frequently in the elderly and in patients with neurodegenerative diseases such as Alzheimer's disease (Maciver, S.K. and Harrington, C.R. (1995) Neuroreport.
30 6:1985-1988). Actinin-4, an actin-bundling protein, appears to be associated with the cell motility of metastatic cancer cells. Other disease associations include premature

chromosome condensation, which is frequently observed in dividing cells from tumor tissue (Murnane, J.P.(1995) Cancer Metastasis Rev. 14:17 29), and the significant roles of axonernal and assembly MAPs in viral pathogenesis (Sodeik, B. et al. (1997) J. Cell Biol. 136:1007 1021).

- 5 Table 1 lists several of the peptides described herein that appear, based upon structural homology, to be cytoskeletal proteins. These peptides (and their corresponding source proteins) can therefore be used to treat disorders associated with inappropriate cytoskeletal protein expression or activity. Examples of such disorders include cell proliferative, immunological, vesicle trafficking, reproductive, smooth muscle, 10 developmental, and nervous disorders, or to screen for agonists and antagonists useful for the same purpose. In addition, nucleic acids encoding the proteins as well as compounds (e.g., antibodies) that recognize the proteins can be used in a wide variety of applications described herein, including therapeutics, diagnostics, and drug screening.

15 The invention therefore includes the following peptides as cytoskeletal proteins:
SEQ ID NO:118, SEQ ID NO:144, SEQ ID NO:177, SEQ ID NO:183, and SEQ ID NO:185.

Receptors

- As described in Examples 1 and 2 (and the accompanying table), many of the 20 peptides described herein are derived from proteins that appear to be receptors. Receptors are a broad category of proteins that specifically recognize other molecules. Many receptors are cell surface proteins that bind extracellular ligands and produce cellular responses in the areas of growth, differentiation, endocytosis, and immune response. Other receptors facilitate the selective transport of proteins out of the 25 endoplasmic reticulum and localize enzymes to particular locations in the cell. The propagation of cellular signals and the transport and localization of proteins rely upon specific interactions between receptors and a variety of associated proteins. Examples of families of receptors include: G-protein Coupled Receptors (GPCRs); MHC molecules; hormone receptors; and TNF receptor superfamily members.
- 30 Receptor-mediated signal transduction is the process whereby cells communicate with one another and respond to extracellular signals via a series of biochemical events.

Extracellular signals are transduced through a biochemical cascade that begins with the binding of a signal molecule to a cell membrane receptor. The signal is propagated to effector molecules by intracellular signal transducing proteins and culminates with the activation of an intracellular target molecule. The process of signal transduction
5 regulates a wide variety of cell functions including cell proliferation, cell differentiation, induction of immune responses, and gene transcription.

Table 1 lists several of the peptides described herein that appear, based upon structural homology, to be receptors. These peptides (and their corresponding source proteins) can therefore be used to treat disorders associated with inappropriate receptor
10 expression or activity. Examples of such disorders include immunological disorders (including autoimmune/inflammatory disorders) and cell proliferative disorders (including cancer), or to screen for agonists and antagonists useful for the same purpose. In addition, nucleic acids encoding the proteins as well as compounds (e.g., antibodies) that recognize the proteins can be used in a wide variety of applications described herein,
15 including therapeutics, diagnostics, and screening.

The invention therefore includes the following peptides as receptors: SEQ ID NO:1, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:41, SEQ ID NO:44, SEQ ID NO:45, SEQ ID NO:46, SEQ ID NO:48, SEQ ID NO:49, SEQ ID NO:51, SEQ ID NO:53, SEQ ID NO:54, SEQ ID NO:55, SEQ ID NO:57, SEQ ID NO:58, SEQ ID NO:59, SEQ ID NO:60, SEQ ID NO:61, SEQ ID NO:62, SEQ ID NO:63, SEQ ID NO:64, SEQ ID NO:66, SEQ ID NO:67, SEQ ID NO:69, SEQ ID NO:77, SEQ ID NO:78, SEQ ID NO:79, SEQ ID NO:80, SEQ ID NO:81, SEQ ID NO:82, SEQ ID NO:83, SEQ ID NO:84, SEQ ID NO:86, SEQ ID NO:88, SEQ ID NO:89, SEQ ID NO:90, SEQ ID NO:91, SEQ ID NO:92, SEQ ID NO:94, SEQ ID NO:100, SEQ ID NO:104, SEQ ID NO:112, SEQ ID NO:121, SEQ ID NO:122, SEQ ID NO:123, SEQ ID NO:128, SEQ ID NO:129, SEQ ID NO:133, SEQ ID NO:135, SEQ ID NO:143, SEQ ID NO:145, SEQ ID NO:147, SEQ ID NO:150, SEQ ID NO:161, SEQ ID NO:164, SEQ ID NO:166, SEQ ID NO:168, SEQ ID NO:170, SEQ ID NO:171, SEQ ID NO:172, SEQ ID NO:173, SEQ ID NO:174, SEQ ID NO:175, SEQ ID NO:176, SEQ ID NO:180, SEQ ID NO:187, SEQ ID NO:188, SEQ ID NO:194, SEQ ID NO:211, SEQ ID NO:217, SEQ ID NO:218, SEQ ID NO:219, SEQ ID NO:221, and SEQ ID NO:230.

Transcription Factors

- As described in Examples 1 and 2 (and the accompanying table), many of the peptides described herein are derived from proteins that appear to be transcription factors.
- 5 Regulation of gene transcription is the primary process by which a cell controls the appropriate expression of the multitude of genes necessary for growth and differentiation. The selective expression of genes at appropriate times is highly specialized in cells of multicellular organisms and permits the cells to perform "housekeeping" functions and respond to changes in their environment. These changes involve extracellular signals
- 10 from a variety of sources such as hormones, neurotransmitters, and growth and differentiation factors.

Gene transcription is controlled by proteins termed transcription factors. Transcription factors act by binding to a short segment of DNA located near the site of transcription initiation. Binding of a transcription factor to the target DNA activates

15 transcription of the gene. Transcription factors contain a variety of structural motifs that, alone or in combination with one another, permit them to recognize and bind to the wide variety of target DNA sequences.

One group of transcription factors, the TFIIIA subclass of zinc-finger proteins, is characterized by an amino acid motif (a cysteine followed by two to four amino acids, a

20 cysteine, twelve amino acids, a histidine, three to four amino acids, and a histidine) that interacts with zinc ions. The carboxyl terminus of the TFIIIA proteins has three of these "zinc finger" motifs and specifically binds to DNA fragments containing a CACCC pattern. The amino-terminal portion of the TFIIIA proteins is proline and serine-rich and can function as a transcriptional activator. TFIIIA proteins are often important for the

25 proper differentiation of tissues in which they are expressed.

Table 1 lists several of the peptides described herein that appear, based upon structural homology, to be transcription factors. These peptides (and their corresponding source proteins) can therefore be used to treat disorders associated with inappropriate transcription factor expression or activity, or to screen for agonists and antagonists useful

30 for the same purpose. Examples of such disorders include cancer, arthritis, and developmental disorders. In addition, nucleic acids encoding the proteins as well as

compounds (e.g., antibodies) that recognize the proteins can be used in a wide variety of applications described herein, including therapeutics, diagnostics, and screening.

The invention therefore includes the following peptides as transcription factors:

SEQ ID NO:2, SEQ ID NO:10, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:22, SEQ
5 ID NO:27, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:40, SEQ ID NO:43, SEQ ID
NO:96, SEQ ID NO:102, SEQ ID NO:117, SEQ ID NO:120, SEQ ID NO:138, SEQ ID
NO:177, SEQ ID NO:183, SEQ ID NO:184, and SEQ ID NO:208.

Therapeutics

10 As described above, many of the peptides of SEQ ID NOs:1-235 belong to biological classes of proteins that have been implicated in a wide variety of disease conditions. These biological classes include kinases, phosphatases, receptors, proteases, transcription factors, transporters (such as ion channels), and cytoskeletal proteins. Additional biological classifications of many of the peptides of SEQ ID NOs:1-235 are
15 detailed in the "biological class" column of the Table. Members of these additional classifications have also been characterized as being associated with specific disorders.

In addition to disorders associated with discrete biological classes, many of the peptides of SEQ ID NOs:1-235 were derived from transformed cells and thus may be involved in cellular proliferative and/or differentiative disorders, e.g., cancer. The
20 Examples and associated table describe in detail the specific transformed cell lines with which the individual peptides of the application have been found to be associated. Because these peptides have been found to be translated in transformed cells, they are expected to be useful in therapeutic, diagnostic, and screening applications as described herein. For example, in one embodiment, a compound that modulates (increases or
25 decreases) the expression or activity of a polypeptide containing any of SEQ ID NOs:1-235 can be used to treat or prevent a cellular proliferative and/or differentiative disorder, e.g., a B cell cancer such as myeloma, colon cancer, gastric cancer, adenocarcinoma, sarcoma, melanoma, lymphoma, or leukemia.

In one embodiment, a polypeptide containing any of SEQ ID NOs:1-235 (or a
30 nucleic acid encoding such a polypeptide) can be administered to a subject to treat a disorder. For example, a disorder characterized by insufficient levels of a given

polypeptide, e.g., a phosphatase or an ion channel, can be treated by such a method. In one example, a secreted protein described herein, e.g., a cytokine, is administered to a subject to treat a disorder.

In one embodiment, antagonists or inhibitors of a polypeptide containing any of SEQ ID NOs:1-235 may be administered to a subject to treat or prevent a disorder. In one aspect, antibodies specific for a polypeptide containing any of SEQ ID NOs:1-235 may be used directly as an antagonist, or indirectly as a targeting or delivery mechanism for bringing a pharmaceutical agent to cells or tissue that expresses the polypeptide.

The invention features a method for treating cancer comprising administering to a patient in need of such treatment an amount of a composition comprising a polypeptide as described herein in an amount sufficient to elicit an immunogenic response. Also, the invention features a method for treating a cancer patient, the method comprising administering to the patient an antibody that selectively binds to a peptide as described herein.

In other embodiments, therapeutic proteins, antagonists, antibodies, agonists, antisense sequences or vectors may be administered in combination with other appropriate therapeutic agents. Selection of the appropriate agents for use in combination therapy may be made by one of ordinary skill in the art, according to conventional pharmaceutical principles. The combination of therapeutic agents may act synergistically to effect the treatment or prevention of the various disorders described above. Using this approach, one may be able to achieve therapeutic efficacy with lower dosages of each agent, thus reducing the potential for adverse side effects.

Antagonists or inhibitors of the polypeptides may be produced using methods which are generally known in the art. In particular, purified polypeptides may be used to produce antibodies or to screen libraries of pharmaceutical agents to identify those which specifically bind the polypeptide. Cells expressing a nucleic acid of the invention can be screened against the same libraries to find agents that bind and/or affect the activity of the encoded polypeptide.

An additional embodiment of the invention relates to the administration of a pharmaceutical composition, in conjunction with a pharmaceutically acceptable carrier, for any of the therapeutic effects discussed above. Such pharmaceutical compositions

may consist of a polypeptide containing any of SEQ ID NOs:1-235, antibodies to the polypeptide, mimetics, agonists, antagonists, or inhibitors of the polypeptide. In addition to the active ingredients, these pharmaceutical compositions may contain suitable pharmaceutically-acceptable carriers comprising excipients and auxiliaries which

5 facilitate processing of the active compounds into preparations which can be used pharmaceutically. Further details on techniques for formulation and administration may be found in the latest edition of Remington's Pharmaceutical Sciences (Maack Publishing Co., Easton, Pa.). The compositions may be administered alone or in combination with at least one other agent, such as a stabilizing compound, which may be administered in any

10 sterile, biocompatible pharmaceutical carrier, including, but not limited to, saline, buffered saline, dextrose solution, and water. The compositions may be administered to a patient alone, or in combination with other agents, drugs or hormones.

Diagnostics

15 In another embodiment, compounds (e.g., antibodies) that specifically bind to a polypeptide containing any of SEQ ID NOs:1-235 may be used for the diagnosis of conditions or diseases characterized by expression of the polypeptide, or in assays to monitor patients being treated with the polypeptide, agonists, antagonists or inhibitors. Antibodies useful for diagnostic purposes may be prepared in the same manner as those

20 prepared for therapeutic purposes. Diagnostic assays for a polypeptide containing any of SEQ ID NOs:1-235 include methods that utilize the antibody and a label to detect the polypeptide in human body fluids or extracts of cells or tissues. The antibodies may be used with or without modification, and may be labeled by joining them, either covalently or non-covalently, with a reporter molecule. A wide variety of reporter molecules that are

25 known in the art may be used, several of which are described above.

In another embodiment of the invention, a polynucleotide, e.g., a polynucleotide encoding a polypeptide containing any of SEQ ID NOs:1-235, may be used for diagnostic purposes. The polynucleotides that may be used include oligonucleotides, antisense RNA and DNA molecules, and PNAs. The polynucleotides may be used to detect and

30 quantitate gene expression in biopsied tissues in which expression of a polypeptide described herein may be correlated with disease. The diagnostic assay may be used to

distinguish between the absence, presence, and excess expression of an mRNA encoding a polypeptide containing any of SEQ ID NOs:1-235, and to monitor regulation of mRNA levels during therapeutic intervention.

- A polynucleotide encoding a polypeptide containing any of SEQ ID NOs:1-235
- 5 may be used for the diagnosis of conditions or diseases that are associated with expression of the polypeptide. Examples of such conditions or diseases include cancers such as cancer of the testis, colon, prostate, uterus, cervix, ovary, lung, intestine, liver, breast, skin, heart, brain, stomach, pancreas, and spleen. The polynucleotide encoding the polypeptide may be used in Southern or northern analysis, dot blot, or other
- 10 membrane-based technologies; in PCR technologies; or in dip stick, pin, ELISA or chip assays utilizing fluids or tissues from patient biopsies to detect altered mRNA expression. Such qualitative or quantitative methods are well known in the art.

As the peptides described herein were found to be translated in transformed cells, these peptides can thus function as markers for a transformed cell, e.g., a cancer cell. As

15 such, detection of polypeptides containing these peptides (or nucleic acids encoding the same) are particularly useful in the diagnosis of cellular proliferative and/or differentiative disorders such as cancer.

Screening Assays

- 20 The invention provides methods for identifying modulators, i.e., candidate or test compounds or agents (e.g., proteins, peptides, peptidomimetics, peptoids, small molecules or other drugs) which bind to a polypeptide containing any of SEQ ID NOs:1-235, have a stimulatory or inhibitory effect on, for example, expression or activity of the polypeptide, or have a stimulatory or inhibitory effect on, for example, the expression or
- 25 activity of a substrate of the polypeptide. Compounds thus identified can be used to modulate the activity of target gene products in a therapeutic protocol, to elaborate the biological function of the target gene product, or to identify compounds that disrupt normal target gene interactions.

The compounds that may be screened in accordance with the invention include,

30 but are not limited to peptides, antibodies and fragments thereof, and other organic

compounds that bind to a polypeptide containing any of SEQ ID NOs:1-235 and increase or decrease an activity of the polypeptide.

- Such compounds may include, but are not limited to, peptides such as soluble peptides, including but not limited to members of random peptide libraries (Lam et al., 5 Nature 354:82 [1991]; Houghten et al., Nature 354:84 [1991]) and combinatorial chemistry-derived molecular libraries made of D- and/or L configuration amino acids; phosphopeptides (including but not limited to members of random or partially degenerate, directed phosphopeptide libraries; Songyang et al., Cell 72:767 [1993]); antibodies (including but not limited to polyclonal, monoclonal, humanized, anti-10 idiotypic, chimeric and single chain antibodies; FAb, F(ab')2 and FAb expression library fragments; and epitope-binding fragments thereof); and small organic or inorganic molecules.

Other compounds that can be screened in accordance with the invention include but are not limited to small organic molecules that are able to gain entry into an 15 appropriate cell and affect (1) the expression of the gene encoding a polypeptide containing any of SEQ ID NOs:1-235 or (2) the activity of the polypeptide.

- As used herein "small molecules" include, but are not limited to, peptides, peptidomimetics (e.g., peptoids), amino acids, amino acid analogs, polynucleotides, 20 polynucleotide analogs, nucleotides, nucleotide analogs, organic or inorganic compounds (i.e., including heteroorganic and organometallic compounds) having a molecular weight less than about 10,000 grams per mole, organic or inorganic compounds having a molecular weight less than about 5,000 grams per mole, organic or inorganic compounds having a molecular weight less than about 1,000 grams per mole, organic or inorganic compounds having a molecular weight less than about 500 grams per mole, and salts, 25 esters, and other pharmaceutically acceptable forms of such compounds.

- Computer modeling and searching technologies permit identification of compounds, or the improvement of already identified compounds, that can modulate expression or activity of a polypeptide containing any of SEQ ID NOs:1-235. Having identified such a compound or composition, the active sites or regions are identified. 30 Such active sites might typically be a binding for a natural modulator of activity. The active site can be identified using methods known in the art including, for example, from

the amino acid sequences of peptides, from the nucleotide sequences of nucleic acids, or from study of complexes of the relevant compound or composition with its natural ligand. In the latter case, chemical or X-ray crystallographic methods can be used to find the active site by finding where on the factor the modulator (or ligand) is found.

5 Although described above with reference to design and generation of compounds that could alter binding, one could also screen libraries of known compounds, including natural products or synthetic chemicals, and biologically active materials, including proteins, for compounds which bind to a polypeptide containing any of SEQ ID NOS:1-235.

10 In vitro systems may be designed to identify compounds capable of interacting with a polypeptide containing any of SEQ ID NOS:1-235. Compounds identified may be useful, for example, in the treatment of conditions such cellular proliferative and differentiative disorders, e.g., cancer.

15 The principle of the assays used to identify compounds that bind to a polypeptide containing any of SEQ ID NOS:1-235 involves preparing a reaction mixture of the polypeptide (or a domain thereof) and the test compound under conditions and for a time sufficient to allow the two components to interact and bind, thus forming a complex which can be removed and/or detected in the reaction mixture. The polypeptide species used can vary depending upon the goal of the screening assay. In some situations it is
20 preferable to employ a peptide corresponding to a domain of the polypeptide fused to a heterologous protein or polypeptide that affords advantages in the assay system (e.g., labeling, isolation of the resulting complex, etc.) can be utilized.

25 The screening assays can be conducted in a variety of ways. For example, one method to conduct such an assay involves anchoring a peptide (or polypeptide or fusion protein) or the test substance onto a solid phase and detecting peptide/test compound complexes anchored on the solid phase at the end of the reaction. In one embodiment of such a method, the peptide reactant may be anchored onto a solid surface, and the test compound, which is not anchored, may be labeled, either directly or indirectly. The invention features a peptide array comprising at least 100 peptides selected from the
30 group consisting of peptides as described herein, each peptide linked to a solid support at a known location. Additionally, the invention features a collection of at least 10

polypeptide arrays, each array comprising at least 100 polypeptides as described herein, each peptide linked to a solid support at a known location. Peptide arrays and methods for producing such arrays are described in, e.g., U.S. Patent No. 5,591,646.

In practice, microtiter plates may conveniently be utilized as the solid phase. The 5 anchored component may be immobilized by non-covalent or covalent attachments. Non-covalent attachment may be accomplished by simply coating the solid surface with a solution of the protein and drying. Alternatively, an immobilized antibody, preferably a monoclonal antibody, specific for the protein to be immobilized may be used to anchor the protein to the solid surface. The surfaces may be prepared in advance and stored.

- 10 In order to conduct the assay, the nonimmobilized component is added to the coated surface containing the anchored component. After the reaction is complete, unreacted components are removed (e.g., by washing) under conditions such that any complexes formed will remain immobilized on the solid surface. The detection of complexes anchored on the solid surface can be accomplished in a number of ways.
- 15 Where the previously non-immobilized component is pre-labeled, the detection of label immobilized on the surface indicates that complexes were formed. Where the previously non-immobilized component is not pre-labeled, an indirect label can be used to detect complexes anchored on the surface; e.g., using a labeled antibody specific for the previously non-immobilized component (the antibody, in turn, may be directly labeled or 20 indirectly labeled with a labeled anti-Ig antibody).

Alternatively, a reaction can be conducted in a liquid phase, the reaction products separated from unreacted components, and complexes detected, e.g., using an immobilized antibody specific for a polypeptide of the invention or the test compound to anchor any complexes formed in solution, and a labeled antibody specific for the other 25 component of the possible complex to detect anchored complexes.

Alternatively, cell-based assays can be used to identify compounds that interact with a polypeptide containing any of SEQ ID NOs:1-235. To this end, cell lines that express the polypeptide, or cell lines that have been genetically engineered to express the polypeptide can be used. Cell based assays are particularly useful for evaluating the 30 functional effects of a compound identified by a screen described herein. For example, once a compound is identified based upon its ability to bind to a polypeptide of the

invention, the compound can then be tested for its ability to, e.g., bind to and/or induce the selective killing of transformed cells.

Use of Peptides and Nucleic Acids Encoding Peptides to Inhibit an Immune Response

- 5 The MHC-binding peptides of SEQ ID NOs:1-235 and the nucleic acids encoding them can be used to block MHC class I and class II-mediated antigen presentation to T cells and thereby inhibit an immune response. Inhibiting an immune response can be particularly useful in conditions such as autoimmune disorders. Methods of using “blocking peptides” to prevent MHC-mediated presentation of antigens to T cells are
- 10 described in U.S. Patent No. 5,827,516. For a polypeptide, e.g., a fusion protein, containing an MHC-binding peptide sequence of any of SEQ ID NOs:1-235, introduction of the polypeptide (or a nucleic acid encoding the polypeptide) to a cell is expected to result in the processing and presentation of the peptide sequence in the context of an MHC class I or class II molecule.
- 15 Peptides described herein may be also useful for inhibiting an immune response when complexed with an MHC molecule, e.g., an HLA molecule, and administered to a host, e.g., a human. The use of HLA/peptide complexes to induce T cell nonresponsiveness has been described for the treatment of autoimmune conditions (see, e.g., Nag et al., 1996, Cell. Immunol. 170:25; Arimilli et al., 1996, Immunol. Cell. Biol. 74:96; Prokaeva, 2000, Curr. Opin. Investig. Drugs 1:70). In addition, antibodies directed against HLA/peptide complexes may be useful in treating disease and/or blocking T cell activation.

Use of Peptides and Nucleic Acids Encoding Peptides as References for MHC Class I and Class II Binding

- 25 Some of the peptides of SEQ ID NOs:1-235 have been characterized as binding to MHC class I or class II molecules (see Example section). These peptides, polypeptides containing them and nucleic acids encoding the same are therefore useful as references in evaluating the ability of a test peptide to bind to an MHC molecule. For example, a
- 30 peptide described herein (a “reference peptide”) can be used in a competitive assay wherein a test peptide is evaluated for its ability to compete with the reference peptide for

binding to an MHC molecule. The reference peptide can optionally be labeled, e.g., with a radioactive label, and displacement of bound label in the presence of a test peptide can be measured. Alternatively, the test peptide can be labeled. Competitive peptide binding assays using a reference peptide are described in, e.g., U.S. Patent 6,037,135.

5

Delivery Systems

- The purified polypeptides, or complexes containing them (such as heat shock protein or MHC complexes), or isolated nucleic acids, can be administered using standard methods, e.g., those described in Donnelly et al. (1994) J. Imm. Methods 176:145, and
- 10 Vitiello et al. (1995) J. Clin. Invest. 95:341. Purified polypeptides and/or isolated nucleic acids of the invention can be injected into subjects in any manner known in the art, e.g., intramuscularly, intravenously, intraarterially, intradermally, intraperitoneally, intravaginally, or subcutaneously, or they can be introduced into the gastrointestinal tract or the respiratory tract, e.g., by inhalation of a solution or powder containing the
- 15 polypeptides or nucleic acids. Alternatively, the purified polypeptides or isolated nucleic acids of the invention may be applied to the skin, or electroporated into the cells or tissue. Purified polypeptides or isolated nucleic acids of the invention may be electroporated with the delivery systems (e.g. microparticles, hydrogels and polymer networks) described herein.
- 20 The purified polypeptides and isolated nucleic acids encoding polypeptides can be delivered in a pharmaceutically acceptable carrier such as saline, lipids, depot systems, hydrogels, networks, liposomes, particulates, virus-like particles, microspheres, or nanospheres; as colloidal suspensions; or as powders. The nucleic acid can be naked or associated or complexed with a delivery vehicle. For a description of the use of naked
- 25 DNA, see, e.g., U.S. Patent No. 5,693,622. For a description of the use of encapsulated DNA see, e.g., U.S. Patent No. 5,783,567. For a description of the use of hydrogel and network delivery systems for DNA delivery see, e.g., USSN 60/262,219. Nucleic acids and polypeptides can be delivered using delivery vehicles known in the art, such as lipids, liposomes, ISCOMS, microspheres, microcapsules, microparticles, gold particles, virus-like particles, nanoparticles, hydrogels or networks, polymers, condensing agents,
- 30 polysaccharides, polyamino acids, dendrimers, saponins, adsorption enhancing materials,

or fatty acids. Viral particles can also be used, e.g., retroviruses, adenovirus, baculovirus, adeno-associated virus, pox viruses, SV40 virus, alpha virus or herpes viruses.

It is expected that a dosage of approximately 0.1 to 100 μ moles of the polypeptide, or of about 1 to 200 μ g of DNA, would be administered per kg of body weight per dose. As is well known in the medical arts, dosage for any given patient depends upon many factors, including the patient's size, body surface area, age, the particular compound to be administered, sex, time and route of administration, general health, and other drugs being administered concurrently. Determination of optimal dosage is well within the abilities of a pharmacologist of ordinary skill.

Other standard delivery methods, e.g., biolistic transfer, or *ex vivo* treatment, can also be used. In *ex vivo* treatment, cells, e.g., antigen presenting cells (APCs), dendritic cells, peripheral blood mononuclear cells, or bone marrow cells, can be obtained from a patient or an appropriate donor and treated *ex vivo* with a composition of the invention, and then returned to the patient.

Microparticles, including those described in U. S. Patent No. 5,783,567 and USSN 60/208,830, can be used as vehicles for delivering macromolecules such as DNA, RNA, or polypeptides into cells. Microparticles may also be made, for example, according to the methods of Mathiowitz, et al. as described in WO 95/24929, herein incorporated by reference. The microparticles can contain macromolecules embedded in a polymeric matrix or enclosed in a shell of polymer. Microparticles act to maintain the integrity of the macromolecule, e.g., by maintaining the DNA in a nondegraded state. Microparticles can also be used for pulsed delivery of the macromolecule, and for delivery at a specific site or to a specific cell or target cell population.

The polymeric matrix can be a synthetic or natural biodegradable co-polymer such as poly-lactic-*co*-glycolic acid, starch, gelatin, or chitin. Microparticles that are less than 10 μ M in diameter can be used in particular to maximize delivery of DNA molecules into a subject's phagocytotic cells. Alternatively, microparticles that are greater than 10 μ M in diameter can be injected or implanted in a tissue, where they form a deposit. As the deposit breaks down, the nucleic acid or polypeptide is released gradually over time and taken up by neighboring cells.

The purified polypeptides and isolated nucleic acids of the invention can be administered by using Immune Stimulating Complexes (ISCOMS), which are negatively charged, cage-like structures of 30-40nm in size formed spontaneously on mixing cholesterol and Quil A (saponin), or saponin alone. A polypeptide (or analog) and nucleic acid of the invention can be co-administered with an ISCOM, or the polypeptide (or analog) and nucleic acid can be administered separately. The polypeptides and nucleic acids of the invention may also be electroporated into cells or tissues of a recipient. Electroporation may occur *ex vivo* or *in vivo*.

10 Peptide Profiles and Databases

U.S. Patent Application 09/372,380, the content of which is herein incorporated by reference, provides compositions and methods for the characterization of a cell's protein repertoire and the storage and manipulation of that information in a computer database. A characteristic profile or fingerprint of peptides or polypeptide ligands can be generated, for example, for a given cell type, for diseased vs. normal cells, and for different metabolic or developmental states of a cell. Appropriate comparisons of the profiles can be used to identify cellular targets useful in diagnostics, drug screening and development, and delivering therapeutic regimens. The EPTs described herein, the MHC-binding peptides of SEQ ID NOs:1-235, represent a population of polypeptide ligands that can be used in the methods, ligand profiles, and databases described in USSN 09/372,380. In addition to EPTs, all of the peptides described herein can be used to catalogue and profile the protein composition of a cell. The following are several non-limiting examples of uses of the peptides for identifying, cataloguing and profiling the protein composition of a cell.

25 Peptides and proteins from which they are derived can be used to identify, catalogue and characterize most or all proteins expressed within a cell for any given cell type, metabolic or developmental stage, and disease vs. normal state, or in response to a test substance such as a given hormone, growth factor, transcription factor, cytokine, small molecule, polypeptide, nucleic acid, carbohydrate or lipid. The approach can also 30 identify differences between transgenic vs. non-transgenic cells, or transfected vs. non-transfected cells. As such, the invention relates to the identification of "polypeptide or

peptide profiles" of a cell type of interest. These profiles can be used to pre-sort cellular proteins for "proteomics" analysis, greatly reducing the screening effort and increasing the efficiency of identifying cellular proteins involved in developmental and metabolic disease processes. Appropriate comparisons of the profiles can be used to identify

- 5 cellular targets useful in diagnostics, drug screening and development, and for developing therapeutic regimens. Such data will facilitate the identification of proteins that have biological significance to a particular cellular state, *e.g.*, in metabolism, maturation, development, disease or treatment.

Peptide esterification methods such as those described in U.S. Provisional Application No. 60/284,416, filed April 16, 2001, the content of which is herein incorporated by reference, can be used to determine relative protein quantities in different cells or tissues.

Peptides of the invention can be used for comparative purposes. A distinct peptide profile, *e.g.*, an EPT profile, can be generated for each cell of interest. The 15 profiles of different cells, tissue or organ types of interest may be compared, and polypeptides may be identified that are differentially represented, *e.g.*, present in one type of cell/tissue/organ, but absent from another, or expressed with different abundance. Furthermore, "differential profiles" of polypeptides may be generated representing peptides that are differentially present in the two types of cells.

Peptides described herein can be used to verify or confirm the distinct profile of a 20 cell of interest. In this use, polypeptides from cells that are essentially identical are isolated and compared. Comparison of the peptide profiles confirms that they are essentially identical, and together represent a reproducible ligand profile for the given cell type. For example, information can be obtained if the peptide profile or set of 25 profiles that represents polypeptides derived from two or more types of MHC molecules in the given cell type are compared. For example, a subtraction profile of polypeptides is generated from comparing polypeptides isolated from two or more types of MHC molecules.

A first cell sample and a second cell sample of interest may be obtained from 30 different types of biological tissue (*e.g.*, comparing smooth muscle tissue to skeletal muscle tissue), different cell types (*e.g.*, endothelial cells and epithelial cells), different

organ systems (e.g., pancreas and lung), or the same organ system but cells of different status (e.g., terminally differentiated vs. embryonic, or healthy vs. diseased or predisposed to a disease). Alternatively, one can compare transfected cells which express a particular recombinant nucleic acid versus non-transfected cells or transfected cells
5 which do not currently express the recombinant nucleic acid. One could also compare cells treated in a particular way (either *in vivo* or *in vitro*) vs. cells treated in a different way, or untreated.

For example, a treatment may involve administration of a test substance or drug candidate such as a growth factor, a hormone, a cytokine, a small molecule, a
10 polypeptide, a nucleic acid, a carbohydrate, or a lipid. Alternatively, a treatment may involve exposing the cells to stress conditions such as trauma, hypoxia, deprivation of glucose, deprivation of an amino acid, deprivation of a nutrient, presence of a toxin, or low or high temperature. The cells are preferably vertebrate cells (e.g., from a bird or fish), and more preferably mammalian cells, e.g., from a human or from a non-human
15 animal such as a non-human primate, a mouse, rat, guinea pig, hamster, rabbit, dog, cat, cow, horse, pig, sheep, or goat. By using a third cell sample, one could compare three different cell samples, or compare the first sample to the second and to the third. For example, the second cell sample could be a positive control and the third cell sample a negative control, or the three cell samples could represent three different treatment
20 regimens.

In a variation on the above, one can simply compare the proteins expressed in a first cell sample to those expressed in a reference cell sample, by generating a peptide profile, e.g., an EPT profile, that is compared to an appropriate reference peptide profile. One compares first peptide profile to the reference peptide profile, in order to identify
25 differences or similarities between the first cell sample and the reference cell sample. This and the other comparison methods described above can be used to compare, for example, cells cultured in the presence of a test compound to cells not cultured in the presence of the test compound; or cells from an animal treated with a test compound to cells (1) from the same animal before the treatment, or (2) from a second animal not treated.

30 Differential peptide profiles can be generated for cells of interest where one peptide profile consists of a subset of polypeptides that is differentially present in two (or

more) distinct cell types, disease stages, developmental stages, metabolic stages, cell cycle stages, treatment regimens, etc., of interest. As such, the differential profiles represent a repertoire of peptides that may directly or indirectly be involved in the different cellular phenotypes or behavior. Consequently, the differential profiles provide

5 a valuable tool for the characterization of cell-type and/or phenotype-specific protein expression, and for the identification and/or the isolation of known or novel gene products and their respective coding sequences that are potentially involved in biological processes, such as developmental processes, establishment and progression of disease, predisposition to disease, organ development, signal transduction, differentiation,

10 neurogenesis, etc., or in response to environmental factors or treatments. For example, the polypeptides identified as differentially expressed may be further characterized by determination of their chemical structure: i.e., sequence. Thus, the present invention provides for the characterization of differential expression, e.g., the presence or absence, of gene products encoded by known genes and/or ESTs with unknown function. The present

15 invention thus can be used as an easy and efficient way to assign to previously identified genes or gene products a putative function and/or involvement or association with a particular developmental pathway, metabolic pathway, or disease stage. With this information, new targets for the development of gene therapy approaches and drug development may rapidly be identified.

20 Peptide profiles for a given cell, tissue or organ of interest can be generated and stored in a database. The compilation of data can then be used for a number of applications. First, they are used as a reference point for a human patient's or animal's sample for the diagnosis of disease, progression of disease, and predisposition for disease. For example, if a disease is associated with changes in protein composition in certain cells, organ systems, cell sources, or tissue types, a suitable patient sample may be used to generate a protein profile, and compared with profiles of corresponding samples of normal (non-diseased) and/or diseased origin to assess presence or absence of, progression of, and/or predisposition to the particular disease in question. A large number of diseases may be diagnosed this way, including diseases for which particular aberrations in protein expression are known,

25 including, but not limited to metabolic diseases that are associated with lack of certain enzymes, proliferative diseases that are associated with aberrant expression of, e.g.,

30

oncogenes or tumor suppressors, developmental diseases that are associated with aberrant gene expression, etc. Furthermore, the peptide profiles can be used for the diagnosis of diseases or other aberrations based on pre-determined differences in EPT profiles. Thus, if it is pre-determined that a given disease of interest is associated with certain changes of the 5 peptide profile of a particular type of cell, tissue, cell source, or organ system, a human patient or animal may be diagnosed based simply on its individual profile when compared to the profiles provided by a database.

Second, peptide information can be used to detect protein translation cell, cell sample, or tissue sample. Such techniques can complement the detection of mRNA and be 10 used to detect specific protein translation (particularly in diseased tissues).

Third, the information stored in a database may be used to identify genes and their products that are involved in the manifestation of, progression of, or predisposition to any disease of interest, and with the development of symptoms of a particular disease. For example, peptide profiles of a diseased organ, tissue or cell type may be generated and 15 compared with the corresponding profile counterpart obtained from a non-diseased sample. Differences in the profile may be identified, and individual peptides that are differentially present in the diseased vs. the non-diseased sample may be identified and isolated for further analysis. The identified differences in the peptide profiles are useful for future diagnosis of the disease or aberration.

20

Generating Peptide Profiles for Different Developmental, Metabolic or Disease Stages of a Given Type of Cell

Peptide profiles for cells of different developmental, metabolic or disease stages can be generated and compared to identify differences in protein or gene expression. For 25 example, the profiles of a cancer cell and non-cancerous cell derived from the same genetically matched tissue may be generated and compared. Proteins differentially expressed in diseased and non-diseased cells can conveniently be identified, and their involvement in disease development and progression analyzed by methods well known in the art. In this way, new targets for the treatment of the disease are efficiently identified.

30 Alternatively, peptide profiles of cells of different developmental stages can be generated and compared. For example, profiles of embryonic cells and adult cells derived

from genetically matched tissue may be generated and compared to identify genes and their products that play a role in developmental processes, and that may be useful for the development of, e.g., novel gene therapy or other therapeutic approaches for the treatment of developmental disorders.

- 5 In another example, peptide profiles of (a) cells infected with a selected pathogen, e.g., microorganism, virus, retrovirus, or prion, and (b) corresponding non-infected cells are generated and compared to identify genes and gene products that are turned on or off in response to the infection. Alternatively, instead of being infected, the first cell can be made to take up a foreign protein or immunogenic substance, etc. This approach allows one, e.g.,
10 to identify factors produced by the cells in response to infection or introduction of the foreign substance that could be useful for therapeutic purposes.

In another example, peptide profiles from cells derived from individuals having a selected genetic disorder and individuals that do not have such disorder are generated and compared. Preferably, samples from affected and non-affected family members are used
15 for the generation of the profiles. Depending on the particular genetic disorder chosen, cell or tissue types that are known to be affected by the particular genetic disorder are studied. In many cases, profiles of various cell and/or tissue types will be generated and compared. This example allows one to identify genes and proteins associated with a genetic disorder. The information obtained may be useful for the development of gene
20 therapy and other therapeutic approaches and for the development of targeted drugs that interfere with the expression of genes or activity or stability of gene products that are involved in the symptoms of the genetic disease. Furthermore, this example allows selection of diagnostic targets for the identification of individuals predisposed for certain types of disease or disease symptoms.

25

Generation of Peptide Profiles Correlated to Response of a Given Cell Type to External Factors

- In one example, a peptide profile of a given cell type treated with an external factor is generated and compared to a profile of cells of the same type which have not
30 been so treated, to identify differences in protein expression. The cells can be

recombinant or native, a cell line or non-transformed cells, or isolated directly from an animal before and after treatment of the animal with the compound.

- For example, peptide profiles of cells of a selected origin or nature that have been contacted with a growth factor, cytokine or hormone, and cells that have not been
- 5 contacted with the substance, but otherwise treated the same way, are generated and compared. This allows identification of genes and gene products that are turned on or turned off in response to the growth factor, cytokine or hormone, which will give, *e.g.*, valuable insight in cellular signal transduction pathways and regulation of protein expression.
- 10 Similarly, peptide profiles of cells that have been treated with or exposed to a polypeptide, small molecule, chemokine, or nucleic acid drug or drug candidate, and cells that have not been treated with or exposed to the substance, but have otherwise been treated the same way, are generated and compared. This allows one to identify the effects of the selected substance on protein expression in the cell, and is, for example, an
- 15 excellent tool for the validation of particular drugs or the identification of drugs associated with expression of a selected gene or gene product.

In another example, peptide profiles of cells that have been exposed to a selected type of compound, *e.g.*, a selected carbohydrate or group of carbohydrates, lipid or group of lipids, amino acid or group of amino acids, nucleotide or nucleoside or group of either, or

20 vitamin or group of vitamins, and cells that have not been treated with the compound, but have otherwise been treated the same way, are generated and compared. This allows one to identify the effects of the selected compound on the gene and protein expression of the cell, and will give valuable insight into metabolic processes.

In another example, peptide profiles of cells that have been treated with a selected nucleic acid, *e.g.*, a selected antisense oligonucleotide, a ribozyme, an expression vector, a plasmid, an RNA, or a DNA, and cells that have not been treated with the nucleic acid, but have otherwise been treated the same way, are generated and compared. This allows one to identify the effects of the antisense oligonucleotide or other nucleic acid on the protein expression in the cell, and as such allows one to evaluate the efficacy or effect of the

25 antisense oligonucleotide or nucleic acid. Antisense molecules and ribozymes of the invention may be prepared by any method known in the art for the synthesis of nucleic

acid molecules. These include techniques for chemically synthesizing oligonucleotides such as solid phase phosphoramidite chemical synthesis. Alternatively, RNA molecules may be generated by *in vitro* and *in vivo* transcription of DNA sequences encoding polypeptides. Such DNA sequences may be incorporated into a wide variety of vectors with suitable RNA polymerase promoters such as T7 or SP6. Alternatively, these cDNA constructs that synthesize antisense RNA constitutively or inducibly can be introduced 5 into cell lines, cells, or tissues.

Finally, peptide profiles of cells that have been subject to a selected stress condition, such as low or high temperature, hypoxia, oxidative stress, free radical-induced stress, deprivation of nutrients such as glucose, amino acids, or other essential factors, or presence of a toxin, are generated and compared to a peptide profile generated in untreated controls. Differentially expressed gene products are identified in order to give valuable insight into factors involved in cellular stress responses. This example provides an extremely valuable and efficient way to determine and/or evaluate the effect 10 of a selected compound on protein expression in the cell. The technique may furthermore be useful to verify a desired shut-down of certain enzymatic activities, *e.g.*, by distinguishing between phosphorylated and non-phosphorylated, or glycosylated and non-glycosylated, peptides and/or proteins. It can also be used to aid in pharmacological 15 and/or toxicological assessment of potential new drugs, and in screening for such drugs.

20

Generating Peptide Profiles for Different Organ Systems

Peptide profiles of cells derived from different organs or organ systems may be generated and compared to identify differences in protein or gene expression. For example, EPT profiles of cells derived from lung, liver, heart, spleen, skin, brain, kidney, 25 thymus, intestine, and/or colon can be generated and compared. Differentially expressed genes and proteins are thus identified. This example is useful to identify proteins that are involved in an organ's particular physiological function.

In another example, peptide profiles of selected tissue or cell types, *e.g.*, muscle, endothelium, epithelium, neuronal, fat, ovarian, testicular, blood, bone marrow, and/or 30 mammary tissue, etc., are generated, compared, and differentially expressed proteins

identified. This will give valuable insight into a protein's involvement in a tissue or cell type's physiological function.

Generating Peptide Profiles for Expression Studies in Standard Cell Lines

- 5 Peptide profiles of cells derived from differentially engineered standard cell lines can be generated and compared to identify differences in protein expression. For example, peptide profiles of standard cell lines that have been engineered to express/overexpress one or several selected recombinant genes, *e.g.*, genes encoding a selected growth factor receptor or other signal transduction component, transcription factor,
- 10 oncogene, apoptosis-inducing gene, etc., are generated and compared to peptide profiles prepared from a reference cell line of the same origin, but which does not carry and express the selected recombinant gene. Differentially expressed genes and gene products are identified. This will allow one to identify the impact of the overexpressed gene on the expression of other polypeptides in the cell.
- 15 The following examples are not to be construed as limiting the scope of the invention in any way.

EXAMPLES

20 Example 1: Isolation and Characterization of MHC Binding Peptides (EPTs)

- This example describes peptides identified by the immunoaffinity purification of class I and class II HLA molecules, followed by acid extraction and solid phase extraction of the EPT repertoire, reversed-phase HPLC separation, and mass spectrometry analysis. Methods used to derive the peptide sequences disclosed in this example are described in detail in U.S. Patent Application 09/372,380, filed August 11, 1999, the content of which is herein incorporated by reference. The various HLA molecules from which peptides were extracted are detailed in Table 1.

- Table 1 describes each of the peptides according to five criteria, as follows: (1) SEQ ID NO; (2) a numeric code corresponding to cell line and HLA type; (3) SEQ ID NOs of source protein reference(s); (4) source protein symbol; and (5) a function key corresponding to biological classification(s).

The SEQ ID NO for each peptide in Table 1 is Criteria 1. The other criteria follow to the right of the peptide sequence and are separated by a vertical hatch divider. Each new peptide entry begins on the next consecutive line having the next consecutive SEQ ID NO.

- 5 Criteria 2 of Table 1 identifies a peptide according to the cell type and HLA type from which it was derived. A numeric code has been assigned to each combination of cell type and HLA type. The numeric code is as follows:

NUMBER	CELL LINE	HLA TYPE
1	721.221	A11
2	721.221	A1
3	721.221	A24
4	721.221	A3
5	721.221	DR1
6	721.221	PAN-DR
7	IM9	A2
8	IM9	N/A
9	JY	A2
10	JY	B7C7
11	JY	DR4_13
12	JY	DR4
13	KATO III	A2
14	KATO III	CLASS 1
15	KATO III	N/A
16	KATO III	PAN-DR
17	LS174T	A2
18	LS180	A2
19	LS180	CLASS 1
20	LS180	PAN-CLASS 2
21	N/A	A11
22	NORMAL PBMC	A2
23	NORMAL PBMC	CLASS 1
24	PRIESS	A2
25	PRIESS	DR4
26	PRIESS	PAN-DR
27	SW403	A2
28	SW480	A2
29	U266	A2

IM-9 is an EBV-transformed B lymphoblastoid cell line derived from the peripheral blood of a patient with multiple myeloma. This cell line is described in, e.g., Fahey et al. (1971) Ann. N.Y. Acad. Sci. 190: 221-234.

U266 is a B lymphocyte cell line established from tissue obtained from a patient with myeloma. This cell line is described in, e.g., Nilsson et al. (1970) Clin. Exp. Immunol. 7:477-489.

LS180 is a human colorectal adenocarcinoma cell line. The cell line is tumorigenic in nude mice. This cell line is described in, e.g., Tom et al. (1976) In Vitro 12:180-191. LS174T is a trypsinized variant of LS180.

SW403 and SW480 are human colorectal adenocarcinoma cell lines. The cell lines are tumorigenic in nude mice. The cell lines are described in, e.g., Fogh et al. (1977) J. Natl. Cancer Inst. 59:221-226.

KATO III is a human gastric cancer cell line. The cell line is described in, e.g., Yamamoto et al. (1996) Cancer 77:1628-33.

JY is a human lymphoblastoid cell line. The cell line is described in, e.g., J. Biol. Chem. (1979) 254:8709, J. Biol. Chem. (1975) 250:4512, and Proc Natl Acad Sci USA (1979) 76:2273.

721.221 is a human lymphoblastoid cell line that has been mutagenized to eliminate the expression of HLA-A, -B, and -C alpha chains. The cell line is described in, e.g., Shimizu et al. (1988) Proc. Natl. Acad. Sci USA 5:227-231. The 721.221 cell lines described herein were transfected with a nucleic acid encoding an individual MHC molecule, e.g., HLA-A1, -A2, -A3, or -A11.

Priess is a human B-lymphoblastoid cell line. The cell line is described in, e.g., Hanania et al. (1983) EMBO J 2:1621-1624.

The SEQ ID NOs of the source protein reference(s) for a given peptide are described as Criteria 3 of Table 1. "Source protein" refers to an amino acid sequence or predicted amino acid sequence contained in a publicly available nucleotide and/or protein database having a region identical to an EPT sequence. In some cases, a "source protein" may not actually represent a protein from which a peptide is derived, but merely a protein (or predicted protein) containing a sequence identical to that of an EPT sequence.

Peptides can be referenced to multiple different source proteins. The list of all identified

source proteins for any one peptide is listed in Table 1. The sequences corresponding to the SEQ ID NOs of the source proteins are in the accompanying sequence listing.

The amino acid sequence for each of the source proteins was derived from NCBI (www.ncbi.nlm.nih.gov/PubMed/). The entire content of this reference is herein
5 incorporated by reference.

- Criteria 4, "source protein symbol," provides the symbol identifying the source protein. Proteins may have been identified by different protein symbols in which case the different protein symbols for the source protein have been listed. Symbols are obtained from three places in the following order: (a) gene symbol(s) and alias(es) from LocusLink; (b) gene name(s) from LocusLink; or (c) Locus titles from LocusLink
10
- Criteria 5, entitled "biological classification," provides a numeric key representing functional classifications for the peptide sequences. Several of these biological classes are described in detail in the application. All known biological classifications for a particular peptide are listed in Table 1. The numeric key
15 corresponding to the biological class is as follows:

FUNCTION	KEY BIOLOGICAL FUNCTION
1	CYTOSKELETON
2	TUMOR SUPPRESSOR
3	DNA BINDING
4	PATHOGENESIS
5	RNA BINDING
6	RIBONUCLEOPROTEIN
7	DNA-BINDING PROTEIN
8	NUCLEUS
9	TRANSCRIPTION CO-REPRESSOR
10	POL II TRANSCRIPTION
11	DNA PACKAGING
12	TRANSFERASE
13	CHROMATIN/CHROMOSOME STRUCTURE
14	TRANSCRIPTION REGULATION
15	HISTONE ACETYLTRANSFERASE
16	TRANSCRIPTION ACTIVATING FACTOR
17	ACTIVATOR
18	INHIBITOR OR REPRESSOR
19	TRANSPORTER
20	POTASSIUM TRANSPORT
21	SULFONYLUREA RECEPTOR
22	CHANNEL [PASSIVE TRANSPORTER]
23	TRANSCRIPTION FACTOR
24	PHOSPHOPYRUVATE HYDRATASE
25	REPRESSION OF TRANSCRIPTION FROM POL II PROMOTER
26	LYASE
27	LEARNING AND MEMORY
28	SYNAPTIC TRANSMISSION
29	SMALL MOLECULE TRANSPORT
30	GLUTAMATE SIGNALLING PATHWAY
31	INTEGRAL PLASMA MEMBRANE PROTEIN
	N-METHYL-D-ASPARTATE SELECTIVE GLUTAMATE
32	RECEPTOR
33	NEURONAL TRANSMISSION
34	RECEPTOR (SIGNALLING)
35	MITOSIS
36	ONCOGENESIS
37	BASEMENT MEMBRANE
38	NUCLEAR CHROMOSOME
39	DNA MEDIATED TRANSFORMATION
40	CHONDROITIN SULFATE PROTEOGLYCAN
41	CHROMOSOME ORGANIZATION AND BIOGENESIS
42	MITOCHONDRION

43	AA3-TYPE CYTOCHROME-C OXIDASE
44	ENERGY GENERATION
45	CYTOSOL
46	5'-NUCLEOTIDASE
47	HYDROLASE
48	METALLOENDOPEPTIDASE
49	PROTEOLYSIS AND PEPTIDOLYSIS
50	NEUROMUSCULAR JUNCTION DEVELOPMENT
51	PROTEASE (OTHER THAN PROTEASOMAL)
52	TRANSCRIPTION CO-FACTOR
53	TRANSCRIPTION FROM POL II PROMOTER
54	RNA POLYMERASE II TRANSCRIPTION FACTOR
55	NEGATIVE CONTROL OF CELL PROLIFERATION
56	CONTROL OF CELL PROLIFERATION
57	CELL PROLIFERATION
58	DEVELOPMENTAL PROCESSES
59	DIFFERENTIATION
60	DNA REPAIR
61	DNA REPLICATION
62	CELL CYCLE CONTROL
63	DELTA-DNA POLYMERASE COFACTOR
64	PROLIFERATING CELL NUCLEAR ANTIGEN
65	DNA SYNTHESIS
66	DNA POLYMERASE OR SUBUNIT
67	CYTOPLASM
68	SOLUBLE FRACTION
69	PROTEIN BIOSYNTHESIS
70	TRYPTOPHANYL-TRNA BIOSYNTHESIS
71	LIGASE
72	PROTEIN SYNTHESIS
73	TRNA SYNTHETASE
74	RNA-BINDING PROTEIN
75	RNA PROCESSING/MODIFICATION
76	LIPID METABOLISM
77	AMINOPHOSPHOLIPID TRANSPORT
78	AMINOPHOSPHOLIPID-TRANSPORTING ATPASE
79	ACTIVE TRANSPORTER, PRIMARY
80	AGEING
81	HELICASE
82	DNA HELICASE
83	3'-5' EXONUCLEASE
84	ADENOSINETRIPHOSPHATASE
85	AGING
86	NUCLEASE [ENDO, EXO, RIBO, DEOXYRIBO]

87	PROTEIN PHOSPHORYLATION
88	NON-MEMBRANE SPANNING PROTEIN TYROSINE KINASE
89	SIGNAL TRANSDUCTION
90	CHROMATIN BINDING
91	EMBRYOGENESIS AND MORPHOGENESIS
92	EMBRYONIC DEVELOPMENT
93	CHROMATIN ARCHITECTURE
94	GLUTAMINE AMIDOTRANSFERASE
95	METABOLISM OF ENERGY RESERVES
96	FRUCTOSE 6-PHOSPHATE METABOLISM
	GLUTAMINE-FRUCTOSE-6-PHOSPHATE TRANSAMINASE
97	(ISOMERIZING)
98	ATP BINDING
99	MITOCHONDRIAL MEMBRANE
100	ATP-BINDING CASSETTE (ABC) TRANSPORTER
101	ATP-BINDING CASSETTE
102	RECEPTOR SIGNALLING PROTEIN
	G-PROTEIN SIGNALLING, LINKED TO CGMP NUCLEOTIDE
103	SECOND MESSENGER
104	PROTEIN KINASE
105	PROTEIN SERINE/THREONINE KINASE
106	NEUROGENESIS
107	NEURONAL DEVELOPMENT
108	PROLINE BIOSYNTHESIS
109	N-ACETYL-GAMMA-GLUTAMYL-PHOSPHATE REDUCTASE
110	OXIDOREDUCTASE
111	AMINO-ACID METABOLISM
112	SNRNP USE
113	MRNA SPLICING
114	MRNA PROCESSING
115	PRE-MRNA SPLICING FACTOR
116	RNA SPLICING
117	SPLICEOSOMAL SUBUNIT
118	TRANSCRIPTION CO-ACTIVATOR
119	REPRODUCTION
120	CELL-TO-CELL SIGNALLING
121	PEROXISOMAL MEMBRANE
122	INTEGRAL PEROXISOMAL MEMBRANE
123	PEROXISOME ORGANIZATION AND BIOGENESIS
124	PEROXISOMAL LONG-CHAIN FATTY ACID IMPORT
125	CNS-SPECIFIC FUNCTIONS
126	IKB KINASE
127	IMMUNE RESPONSE
128	PHOSPHORYLATION OF I-KAPPAB

129	ANTI-PATHOGEN RESPONSE
130	ACTIN BUNDLING
131	ACTIN CYTOSKELETON
132	CELL SHAPE AND CELL SIZE CONTROL
133	ACTIN CYTOSKELETON REORGANIZATION
134	CELL STRUCTURE
135	COMPLEX ASSEMBLY PROTEIN
136	G1/S-SPECIFIC CYCLIN
137	REGULATORY SUBUNIT
138	APOPTOSIS INHIBITOR
139	HISTONE DEACETYLASE
140	ZINC BINDING
141	ANTIMICROBIAL HUMORAL RESPONSE
142	SMALL MOLECULE-BINDING PROTEIN
143	PHOSPHOMANNOMUTASE
144	PROTEIN GLYCOSYLATION
145	N-LINKED GLYCOSYLATION
146	GDP-MANNOSE BIOSYNTHESIS
147	MEMBRANE FRACTION
148	BRAIN DEVELOPMENT
149	PROTEIN BINDING
150	TRANSCRIPTION FACTOR COMPLEX
151	TRANSCRIPTION REGULATION FROM POL II PROMOTER
152	GAS EXCHANGE
153	MICROSOME
154	PLASMA MEMBRANE
155	VESICLE TARGETING
156	ER TO GOLGI TRANSPORT
157	DYSTROPHIN-ASSOCIATED GLYCOPROTEIN COMPLEX
158	ELECTRON TRANSPORTER
159	QUINOLINATE SYNTHASE
160	OTHER METABOLISM
161	CELL MOTILITY
162	INVASIVE GROWTH
163	CYTOSKELETAL STRUCTURAL PROTEIN
164	PERCEPTION OF PEST/PATHOGEN/PARASITE
165	DNA RECOMBINATION
166	PROTEIN MODIFICATION
167	DOUBLE-STRAND BREAK REPAIR
168	SPLICEOSOME
169	ENDONUCLEASE
170	BASE-EXCISION REPAIR
171	ENDODEOXYRIBONUCLEASE
172	URACIL-DNA GLYCOSYLASE

173	DNA-(APURINIC OR APYRIMIDINIC SITE) LYASE
174	CENTROMERE
175	KINETOCHORE
176	ANTI-APOPTOSIS
177	CHROMOSOME SEGREGATION
178	NUCLEAR INNER MEMBRANE, INTEGRAL PROTEIN
179	CELL DEATH/APOPTOSIS
180	ENERGY PATHWAYS
181	GLYCOGEN METABOLISM
182	1,4-ALPHA-GLUCAN BRANCHING ENZYME
183	ENERGY STORAGE
184	CELLULAR DEFENSE RESPONSE
185	CLASS I MAJOR HISTOCOMPATIBILITY COMPLEX ANTIGEN
186	CELL ADHESION
187	ADHESIN/AGGLUTININ
188	NUCLEOLUS
189	NUCLEOPLASM
190	POLY-PYRIMIDINE TRACT BINDING
191	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN
192	DEFENSE RESPONSE
193	LYMPHOCYTE ANTIGEN
194	VISION
195	EXTRACELLULAR MATRIX
196	PHOTORECEPTION
197	EXTRACELLULAR MATRIX COMPONENT
198	PEROXISOME
199	CHROMATIN MODELLING
200	CELL-CELL MATRIX ADHESION
201	CELL MIGRATION/MOTILITY
202	TRANSLATION ELONGATION FACTOR
203	HYDROXYMETHYLGLUTARYL-COA SYNTHASE
204	DNA METHYLATION
205	DNA (CYTOSINE-5-)-METHYLTRANSFERASE
206	STEROL CARRIER
207	STEROL TRANSPORTER
208	ESTRADIOL 17 BETA-DEHYDROGENASE
209	INTEGRAL MEMBRANE PROTEIN
210	ETHANOLAMINEPHOSPHOTRANSFERASE
211	INDUCTION OF APOPTOSIS
212	PROTEIN KINASE CASCADE
213	CHAPERONE
214	CALCIUM BINDING
215	PROTEIN SECRETION
216	ENDOPLASMIC RETICULUM

217	ENDOPLASMIC RETICULUM MEMBRANE
218	CHAPERONES
219	PROTEIN TRANSLOCATION
220	HISTONE DEACETYLASE COMPLEX
221	CELL CYCLE REGULATOR
222	DNA DAMAGE RESPONSE, ACTIVATION OF P53
223	HEAVY METAL BINDING
224	EPIDERMAL DEVELOPMENT AND MAINTENANCE
225	HYDROGEN-TRANSPORTING ATP SYNTHASE
226	HYDROGEN-TRANSPORTING TWO-SECTOR ATPASE
227	CHROMATIN SILENCING
228	ATP DEPENDENT RNA HELICASE
229	RESPONSE TO VIRUSES
230	ANTIVIRAL RESPONSE PROTEIN
231	PATHOGENIC INVASION
232	PEPTIDE TRANSPORT
233	PEPTIDE TRANSPORTER
234	ENZYME ACTIVATOR
235	NON-SELECTIVE VESICLE TRANSPORT
236	TRANSLATION FACTOR
237	PROTEIN SYNTHESIS ELONGATION
238	ACTIN FILAMENT
239	GTPASE ACTIVATOR
240	GTPASE INHIBITOR
241	CALMODULIN BINDING
242	PERIPHERAL PLASMA MEMBRANE PROTEIN
243	GTPASE ACTIVATING PROTEIN
244	LYSOSOME
245	LYSOSOME ORGANIZATION AND BIOGENESIS
246	RNA PROCESSING
247	PEPTIDASE
248	ARSENITE TRANSPORTER
249	NUCLEOCYTOPLASMIC TRANSPORT
250	NUCLEAR-CYTOPLASMIC TRANSPORT
251	GLUCOSE CATABOLISM
252	GLYCEROL-3-PHOSPHATE DEHYDROGENASE
253	NUTRIENT ABSORPTION
254	CYCLOPHILIN
255	ISOMERASE
256	SPECIFIC RNA POLYMERASE II TRANSCRIPTION FACTOR
257	PROTEIN FOLDING
258	HETEROCHROMATIN
259	VIRULENCE
260	H3/H4 HISTONE ACETYLTRANSFERASE

261	METALLOCARBOXYPEPTIDASE
262	CASEIN KINASE II
263	JAK-STAT CASCADE
264	ACUTE-PHASE RESPONSE HEMATOPOEITIN/INTERFERON-CLASS (D200-DOMAIN)
265	CYTOKINE RECEPTOR SIGNAL TRANSDUCER
266	SIALYLTRANSFERASE
267	AMINOSUGAR METABOLISM
268	GLYCOLIPID METABOLISM
269	LIPID:PROTEIN MODIFICATION
270	DNA TOPOISOMERASE
271	DNA TOPOISOMERASE (ATP-HYDROLYZING)
272	TOPOISOMERASE
273	DNA METABOLISM
274	DNA-DIRECTED RNA POLYMERASE I
275	TRANSCRIPTION FROM POL I PROMOTER
276	RNA POLYMERASE I TRANSCRIPTION FACTOR COMPLEX
277	POL I TRANSCRIPTION
278	RNA POLYMERASE SUBUNIT
279	TYROSINE RECOMBINASE
280	26S PROTEASOME
281	19S PROTEASOME REGULATORY PARTICLE
282	PROTEIN DEGRADATION
283	PROTEASOME SUBUNIT
284	ASPARTIC-TYPE ENDOPEPTIDASE
285	GUANYLATE CYCLASE
286	RECEPTOR GUANYLATE CYCLASE
287	MEIOTIC RECOMBINATION
288	MITOTIC RECOMBINATION
289	RNA PROCESSING
290	SMALL NUCLEOLAR RNA
291	OTHER DEVELOPMENT
292	MALE MEIOSIS
293	TRANSCRIPTION FACTOR TFIIE
294	TRANSCRIPTION INITIATION FROM POL II PROMOTER
295	GENERAL RNA POLYMERASE II TRANSCRIPTION FACTOR
296	ARYLESTERASE
297	TUMOR ANTIGEN
298	INFLAMMATORY RESPONSE
299	ANTIBACTERIAL HUMORAL RESPONSE
300	RESPONSE TO PATHOGENIC BACTERIA
301	ATP DEPENDENT DNA HELICASE
	PROTEIN COMPLEX ASSEMBLY, MULTICHAPERONE
302	PATHWAY

303	CARBOHYDRATE METABOLISM
304	ISOCITRATE DEHYDROGENASE (NAD+)
305	GTPASE
306	GTP-BINDING PROTEIN/GTPASE
307	RNA TRANSCRIPTION
308	TRNA TRANSCRIPTION
309	TRANSCRIPTION FACTOR TFIIC
310	TRANSCRIPTION FROM POL III PROMOTER
311	RNA POLYMERASE III TRANSCRIPTION FACTOR
312	POL III TRANSCRIPTION
313	INTRACELLULAR SIGNALLING CASCADE
314	GOLGI APPARATUS
315	ARF GUANYL-NUCLEOTIDE EXCHANGE FACTOR
316	GUANINE NUCLEOTIDE EXCHANGE FACTOR
317	RNA ELONGATION FROM POL II PROMOTER
318	POSITIVE TRANSCRIPTION ELONGATION FACTOR
319	MICROTUBULE
320	STRUCTURAL PROTEIN
321	GLIA CELL DIFFERENTIATION
322	PHOSPHOLIPID BINDING
323	SKELETAL DEVELOPMENT
324	CARTILAGE CONDENSATION
325	BONE DEVELOPMENT AND MAINTENANCE
326	NUCLEAR PORE
327	RAN PROTEIN BINDING
328	IMPORTIN, BETA-SUBUNIT
329	NLS-BEARING SUBSTRATE-NUCLEUS IMPORT
330	NUCLEAR LOCALIZATION SEQUENCE BINDING
331	RECEPTOR (PROTEIN TRANSLOCATION)
332	PROTEIN COMPLEX ASSEMBLY
333	PROLYL-TRNA BIOSYNTHESIS
334	GLUTAMYL-TRNA BIOSYNTHESIS
335	PROTEIN ADP-RIBOSYLATION
336	CELL GROWTH AND MAINTENANCE
337	NAD(+) ADP-RIBOSYLTRANSFERASE
338	LARGE RIBOSOMAL SUBUNIT
339	STRUCTURAL PROTEIN OF RIBOSOME
340	GENERAL CELLULAR ROLE
341	RIBOSOMAL SUBUNIT
342	MEMBRANE.
343	CIRCULATION
344	POSITIVE CONTROL OF CELL PROLIFERATION
345	ANGIOGENESIS
346	HEPARIN N-DEACTYLASE/N-SULFOTRANSFERASE

347	TELOMERE MAINTENANCE
348	REGULATION OF MITOTIC RECOMBINATION
	SINGLE-STRANDED DNA SPECIFIC
349	ENDODEOXYRIBONUCLEASE
350	MAP KINASE
351	TGFBETA RECEPTOR SIGNALLING PATHWAY
352	TRANSLATIONAL REGULATION
353	PROTEIN KINASE INHIBITOR
354	CHEMOTAXIS
355	PHOSPHODIESTERASE I
356	PHOSPHATE METABOLISM
357	NUCLEOTIDE PYROPHOSPHATASE
358	TRANSCRIPTION FACTOR BINDING
	G-PROTEIN LINKED RECEPTOR PROTEIN SIGNALLING
359	PATHWAY
360	OTHER PHOSPHATASE
361	GUANYLATE KINASE
362	OTHER KINASE
363	MOTOR
364	NON-MUSCLE MYOSIN
365	MOTOR PROTEIN
366	DEOXYCYTIDINE KINASE
367	PYRIMIDINE NUCLEOTIDE METABOLISM
368	CHOLINE KINASE
369	LIPID TRANSPORT
370	HEARING
371	CELL CYCLE ARREST
372	MITOTIC G1/S TRANSITION
373	INDUCTION OF APOPTOSIS BY INTRACELLULAR SIGNALS
374	PROTEIN PHOSPHATASE 1 BINDING
375	PROTEIN KINASE A ANCHORING PROTEIN
376	ANCHOR PROTEIN
377	LIPID BINDING
378	ACTIVATION OF MAPK
379	G-PROTEIN LINKED RECEPTOR
380	PHOSPHORYLASE KINASE
381	TRANSCRIPTION TERMINATION FROM POL II PROMOTER
382	CENTRAL NERVOUS SYSTEM DEVELOPMENT
383	OXIDATIVE STRESS RESPONSE
384	CELL STRESS
385	RECEPTOR
386	CELL SURFACE RECEPTOR LINKED SIGNAL TRANSDUCTION
387	LIGAND
388	APOPTOSIS

389	CELL-CELL SIGNALLING
390	HEAT SHOCK PROTEIN
391	TRANSLATIONAL REGULATION, INITIATION
392	MRNA CLEAVAGE
393	MRNA POLYADENYLATION
394	CHOLINESTERASE
395	LAMININ RECEPTOR
396	LAMININ RECEPTOR PROTEIN
397	CYTOSOLIC SMALL RIBOSOMAL (40S)-SUBUNIT
398	FATTY ACID DESATURATION
399	EGF RECEPTOR DOWN REGULATION
400	MICROTUBULE NUCLEATION
401	MICROTUBULE ASSOCIATED PROTEIN
402	INTERLEUKIN-2 RECEPTOR
403	INTERLEUKIN-4 RECEPTOR
404	INTERLEUKIN-7 RECEPTOR
405	INTEGRIN
406	COLLAGEN BINDING
407	BLOOD COAGULATION
408	CELL ADHESION RECEPTOR
409	HISTOGENESIS AND ORGANOGENESIS
410	BLOOD CLOTTING
411	TRANSMEMBRANE RECEPTOR PROTEIN TYROSINE KINASE
412	CYTOSOLIC LARGE RIBOSOMAL (60S)-SUBUNIT
413	CLASS II MAJOR HISTOCOMPATIBILITY COMPLEX ANTIGEN
414	ATP-GATED CATION CHANNEL
415	GONAD DEVELOPMENT
416	GERM CELL MIGRATION
417	CHOLESTEROL METABOLISM
418	CHOLESTEROL BIOSYNTHESIS
419	GERMLINE MAINTENANCE
420	GOLGI CIS-FACE
421	DYNAMIN GTPASE
422	CELL COMMUNICATION
423	MITOCHONDRIAL MEMBRANE ORGANIZATION AND BIOGENESIS
424	1-PHOSPHATIDYLINOSITOL 3-KINASE
425	INOSITOL/PHOSPHATIDYLINOSITOL KINASE
426	PROTEIN-PEROXISOME TARGETING
427	PEROXISOME TARGETING SIGNAL-1 RECEPTOR
428	RAS GTPASE ACTIVATOR
429	CELL CYCLE
430	JNK CASCADE
431	INACTIVATION OF MAPK

432	MESODERM DEVELOPMENT
433	INDUCTION OF APOPTOSIS BY EXTRACELLULAR SIGNALS
434	MISMATCH REPAIR
435	DNA REPAIR ENZYME
436	DNA REPAIR PROTEIN
	EUKARYOTIC TRANSLATION INITIATION FACTOR 2
437	COMPLEX
438	EXTRACELLULAR SPACE
439	BLOOD COAGULATION FACTOR IX
440	PROTEIN PHOSPHATASE TYPE 1 CATALYST
	REGULATION OF G-PROTEIN LINKED RECEPTOR PROTEIN
441	SIGNALLING PATHWAY
442	METHYL TRANSFERASE
443	3'(2'),5'-BISPHOSPHATE NUCLEOTIDASE
	NUCLEOBASE, NUCLEOSIDE, NUCLEOTIDE AND NUCLEIC
444	ACID METABOLISM
445	CYTOCHROME-C OXIDASE
446	LEARNING
447	FEEDING BEHAVIOR
448	PROTEIN TYROSINE KINASE
449	METHIONINE-TRNA LIGASE
450	ACTIN MODULATING
451	NUCLEOTIDE-EXCISION REPAIR
452	SINGLE-STRANDED DNA BINDING
453	EXTRACELLULAR MATRIX STRUCTURAL PROTEIN
454	ADENOSINE DEAMINASE
455	ADENOSINE DEAMINASE REACTION
456	RAS PROTEIN SIGNAL TRANSDUCTION
457	RAL GUANYL-NUCLEOTIDE EXCHANGE FACTOR
458	SMALL GTPASE REGULATORY/INTERACTING PROTEIN
459	COATED VESICLE
460	SECRETORY VESICLE
461	VESICLE TRANSPORT
462	VESICLE COAT PROTEIN
463	HIGH-DENSITY LIPOPROTEIN
464	INTERCELLULAR TRANSPORT
465	DNA DAMAGE RESPONSE
466	EYE PIGMENT BIOSYNTHESIS
467	INTRACELLULAR PROTEIN TRAFFIC
468	PROTEIN DEPHOSPHORYLATION
469	PROTEIN TYROSINE PHOSPHATASE
470	RIBOSOME BIOGENESIS
471	UBIQUITIN LIGASE COMPLEX
472	UBIQUITIN CONJUGATING ENZYME

473	UBIQUITIN-DEPENDENT PROTEIN DEGRADATION
474	PROTEIN CONJUGATION FACTOR
475	CHAPERONIN ATPASE
476	NUCLEIC ACID BINDING
477	HEAT SHOCK RESPONSE
478	NADPH:QUINONE REDUCTASE
479	PHOSPHOGLYCERATE KINASE
480	FK506 BINDING
481	MITOCHONDRIAL MATRIX
482	ELECTRON TRANSFER FLAVOPROTEIN
483	PROTEIN PHOSPHATASE TYPE 1
484	MITOTIC CHECKPOINT
485	ANAPHASE-PROMOTING COMPLEX
486	SIGNAL RECOGNITION PARTICLE
487	DIACYLGLYCEROL KINASE
488	PHOSPHOLIPASE C ACTIVATION
489	CYTOSTOLIC CALCIUM ION CONCENTRATION ELEVATION
490	PHOSPHORIBOSYLGlycinamide FORMYLTRANSFERASE
491	TRANSLATION INITIATION FACTOR EUkARYOTIC TRANSLATION INITIATION FACTOR 3
492	COMPLEX
493	RAN GTPASE ACTIVATOR
494	SIGNAL SEQUENCE RECEPTOR
495	CO-TRANSLATIONAL MEMBRANE TARGETING
496	DEOXYRIBONUCLEOSIDE MONOPHOSPHATE BIOSYNTHESIS
497	IMPORTIN, ALPHA-SUBUNIT
498	REGULATION OF DNA RECOMBINATION
499	NUCLEAR IMPORT/EXPORT PROTEIN
500	SPINDLE
501	CENTROSOME
502	CYTOKINESIS
503	SPINDLE POLE BODY
504	POLYSOME
505	MITOTIC SPINDLE CHECKPOINT CARBAMOYL-PHOSPHATE SYNTHASE (GLUTAMINE-) HYDROLYZING)
507	DEUBIQUITYLATION
508	CYSTEINE-TYPE ENDOPEPTIDASE
509	UBIQUITIN-SPECIFIC PROTEASE
510	ENDOCYTOSIS
511	RAB GTPASE ACTIVATOR
512	INSULIN RECEPTOR SIGNALLING PATHWAY
513	RNA HELICASE
514	LYSINE--tRNA LIGASE

515	NUCLEOSOME ASSEMBLY
516	CHROMATIN ASSEMBLY COMPLEX
517	CALCIUM ION TRANSPORTER
518	INOSITOL-1,4,5-TRIPHOSPHATE RECEPTOR
519	DNA-DIRECTED RNA POLYMERASE II
520	ASPARTATE CATABOLISM
521	CYTOCHROME P450
522	EYE MORPHOGENESIS
523	EXOCYTOSIS
524	SNAP RECEPTOR
525	MEMBRANE FUSION
526	NON-SELECTIVE VESICLE TARGETING
527	DOCKING PROTEIN
528	PROTEIN TARGETING
529	REGULATION OF CDK ACTIVITY
	EUKARYOTIC TRANSLATION INITIATION FACTOR 4
530	COMPLEX
531	SNRNP U2E
532	SNRNP U1E
533	SMALL NUCLEAR RIBONUCLEOPROTEIN
534	PROTEIN LOCALIZATION
535	SERPIN
536	ENZYME INHIBITOR
537	N-METHYLTRANSFERASE
538	N-TERMINAL PROTEIN METHYLATION
539	IMP CYCLOHYDROLASE
	PHOSPHORIBOSYLAMINOIMIDAZOLECARBOXAMIDE
540	FORMYLTRANSFERASE
541	MITOTIC G2 PHASE
	SPINDLE POLE BODY AND MICROTUBULE CYCLE (SENSU 542 SACCHAROMYCES)
543	GMP SYNTHASE
544	PURINE BASE BIOSYNTHESIS
545	DNA DEPENDENT DNA REPLICATION
546	DNA REPLICATION FACTOR A COMPLEX
547	NUCLEOTIDE BINDING
548	DNA REPLICATION CHECKPOINT
549	DNA REPLICATION INHIBITION
550	MITOTIC START CONTROL POINT
551	TEMPERATURE RESPONSE
552	TRANSCRIPTION
553	RECEPTOR SIGNALLING PROTEIN TYROSINE KINASE
554	DAMAGED DNA BINDING
555	PYrimidine-DIMER REPAIR, DNA DAMAGE EXCISION

556	VIRAL REPLICATION
557	PROTEIN C-TERMINUS BINDING
558	PROTEIN PHOSPHATASE TYPE 2A
559	PROTEIN PHOSPHATASE TYPE 2A REGULATOR
560	NUCLEAR MEMBRANE
561	MALATE METABOLISM
562	TRICARBOXYLIC ACID CYCLE MALATE DEHYDROGENASE (OXALOACETATE)
563	DECARBOXYLATING) (NADP+)
564	MEIOSIS
565	SPERMATOGENESIS
566	SISTER CHROMATID COHESION
567	CYCLIN
568	CHOLESTEROL CATABOLISM
569	STEROID HORMONE RECEPTOR
570	TRIPEPTIDYL-PEPTIDASE II
571	D-ALANYL-D-ALANINE ENDOPEPTIDASE
572	PHOSPHOLIPASE C
573	PHOSPHOLIPID METABOLISM
574	TRANSCRIPTION-COUPLED REPAIR
575	ACTIVATION OF JUN KINASE
576	ADENYLYLATE KINASE
577	MICROTUBULE BINDING
578	MITOTIC G2/M TRANSITION
579	STRESS RESPONSE
580	CONTROL OF HEART
581	MUSCLE CONTRACTION
582	STRUCTURAL PROTEIN OF MUSCLE
583	MUSCLE ACTION
584	EPIDERMAL DIFFERENTIATION
585	BILIVERDIN REDUCTASE
586	MAP KINASE
587	VITAMIN METABOLISM
588	RETINOID-X RECEPTOR
589	RETINOIC ACID RECEPTOR
590	IRON HOMEOSTASIS
591	CENTRIOLE
592	CENTROSOME CYCLE
593	TRANSCRIPTION ELONGATION FACTOR
594	RNA DEPENDENT ADENOSINETRIPHOSPHATASE
595	ISOPRENOID BIOSYNTHESIS
596	ISOPENTENYL-DIPHOSPHATE DELTA-ISOMERASE
597	TRANSCRIPTION REGULATION FROM POL III PROMOTER
598	CYCLIN-DEPENDENT PROTEIN KINASE

599	MUSCLE MYOSIN
600	PHOSPHOLIPASE A2
601	MITOTIC CHROMOSOME SEGREGATION
602	OLIGOSACCHARYL TRANSFERASE
603	MYOSIN ATPASE
604	GTP BINDING
605	CELL SHAPE CONTROL
606	SMALL GTPASE MEDIATED SIGNAL TRANSDUCTION
	HYDROGEN-TRANSPORTING ATP SYNTHASE, CATALYTIC
607	CORE
608	MUSCLE DEVELOPMENT
609	PREGNANCY
610	UBIQUITIN--PROTEIN LIGASE
611	ACYLTRANSFERASE
612	FATTY ACID METABOLISM
613	GUANYLATE CYCLASE, SOLUBLE
614	NO MEDIATED SIGNAL TRANSDUCTION
615	FATTY ACID CATABOLISM
616	PROPIONYL-COA CARBOXYLASE
617	PROTEIN ACETYLATION
618	NUCLEOSOME REMODELLING COMPLEX
619	CASPASE-2
620	APOPTOTIC PROGRAM
621	7-ALPHA-HYDROXYSTEROID DEHYDROGENASE
622	INOSITOL-1,4,5-TRIPHOSPHATE 5-PHOSPHATASE
623	LIGAND-DEPENDENT NUCLEAR RECEPTOR
624	ACTIN BINDING
625	POST GOLGI TRANSPORT
626	PYRIDOXAL KINASE
627	LAMIN
628	NUCLEAR LAMINA
629	SERINE PROTEASE INHIBITOR
630	STEAROYL-COA 9-DESATURASE
631	HEPATOCYTE GROWTH FACTOR RECEPTOR
632	BILE ACID BIOSYNTHESIS
633	OXYSTEROL 7-ALPHA-HYDROXYLASE
634	POSTERIOR MIDGUT DEVELOPMENT
635	EXTRACELLULAR
636	MICROTUBULE STABILIZATION
637	EGF RECEPTOR SIGNALLING PATHWAY
638	NEUROTROPHIN TRKA RECEPTOR
	TRANSMEMBRANE RECEPTOR PROTEIN TYROSINE KINASE
639	SIGNALLING PATHWAY
640	METALLOPEPTIDASE

641	MITOCHONDRIAL TRANSLOCATION
642	MITOCHONDRIAL INNER MEMBRANE TRANSLOCASE COMPLEX
643	FRUCTOSE METABOLISM
644	CYTOPLASMIC DYNEIN
645	ARP2/3 PROTEIN COMPLEX
646	CELL ELONGATION
647	NADH DEHYDROGENASE (UBIQUINONE)
648	GAMETOGENESIS
649	MEIOTIC CHROMOSOME
650	DNA DAMAGE CHECKPOINT
651	MITOTIC CHROMOSOME CONDENSATION
652	DNA REPLICATION AND CHROMOSOME CYCLE
653	CAMP-DEPENDENT PROTEIN KINASE REGULATOR
654	EUKARYOTIC TRANSLATION ELONGATION FACTOR 1
655	GOLGI MEMBRANE
656	MANNOSE BINDING LECTIN
657	PHENYLALANINE-TRNA LIGASE
658	PHENYLALANYL-TRNA BIOSYNTHESIS
659	LIGAND BINDING OR CARRIER
660	ELECTRON DONOR
661	ACYL-COA OXIDASE
662	CELL AGEING
663	DNA-DIRECTED RNA POLYMERASE III
664	TRANSCRIPTION REGULATION FROM POL I PROMOTER
665	RIBOSOME
666	SIGNAL RECOGNITION PARTICLE RECEPTOR
667	LONG-CHAIN-FATTY-ACID-COA-LIGASE
668	MONOOXYGENASE
669	TRANSLATIONAL ATTENUATION
670	TROPOMYOSIN BINDING
671	ACTIN CAPPING PROTEIN
672	CHROMATIN
673	PROTEIN-NUCLEUS IMPORT
674	LAMININ-5
675	DEFENSE/IMMUNITY PROTEIN
676	LANOSTEROL 14-ALPHA-DEMETHYLASE
677	SH3/SH2 ADAPTOR PROTEIN
678	RHO PROTEIN SIGNAL TRANSDUCTION
679	ACTIN FILAMENT SEVERING
680	ACTIN POLYMERIZATION/DEPOLYMERIZATION
681	RAB GDP-DISSOCIATION INHIBITOR
682	XENOBIOTIC METABOLISM
683	DETOKSIFICATION RESPONSE

684	CYTOCHROME B5 REDUCTASE
685	NITRIC OXIDE BIOSYNTHESIS
686	NAD(P)H DEHYDROGENASE (QUINONE)
687	SYNAPTIC TRANSMISSION, CHOLINERGIC
688	LAMIN BINDING
689	LAMIN/CHROMATIN BINDING
690	AMYLOID PROTEIN
691	MRNA BINDING
692	GDP-DISSOCIATION INHIBITOR
693	METHENYLtetrahydrofolate cyclohydrolase
694	METHYLENETETRAHYDROFOLATE DEHYDROGENASE
695	SATELLITE DNA BINDING
696	LIPID PARTICLE
	NON-MEMBRANE SPANNING PROTEIN TYROSINE
697	PHOSPHATASE
698	SUPEROXIDE METABOLISM
699	[E]IF-5A]-DEOXYHYPUSINE SYNTHASE
700	COMPLEX I (NADH TO UBIQUINONE)
701	M PHASE
702	CYTOSKELETAL PROTEIN BINDING PROTEIN
703	PHOSPHOLIPASE A1
704	PHOSPHATIDYL SERINE METABOLISM
705	UBIQUITIN ACTIVATING ENZYME
706	SPERMATID DEVELOPMENT
707	DNA REPLICATION ORIGIN BINDING
708	DNA REPLICATION FACTOR
709	DNA REPLICATION FACTOR C COMPLEX
710	MITOTIC G1 PHASE
711	TETRACYCLINE TRANSPORTER
712	ACTIVE TRANSPORTER, SECONDARY
713	MAJOR FACILITATOR SUPERFAMILY
714	PURINE NUCLEOTIDE BIOSYNTHESIS
715	AMIDOPHOSPHORIBOSYLTRANSFERASE
716	PROTEIN-NUCLEUS IMPORT, TRANSLOCATION
717	INTEGRAL PLASMA MEMBRANE PROTEOGLYCAN
718	DNA STRAND ELONGATION
719	TRANSKETOLASE
720	ENDOSOME
721	IRON TRANSPORT
722	TRANSFERRIN RECEPTOR
723	BLOOD PRESSURE REGULATION
724	HETEROTRIMERIC G-PROTEIN GTPASE, BETA SUBUNIT
	ACETYL CHOLINE RECEPTOR SIGNALLING, MUSCARINIC
725	PATHWAY

726	PROTON TRANSPORT
727	VACUOLAR HYDROGEN-TRANSPORTING ATPASE
728	SODIUM/POTASSIUM-EXCHANGING ATPASE
729	SODIUM/POTASSIUM-TRANSPORTING ATPASE
730	ANION TRANSPORT
731	MITOCHONDRIAL OUTER MEMBRANE
732	VOLTAGE-DEPENDENT ANION CHANNEL PORIN
733	APOPTOGENIC CYTOCHROME C RELEASE CHANNEL
734	ADENINE TRANSPORT
735	DOUBLE-STRANDED DNA BINDING
736	CALCIUM-TRANSPORTING ATPASE
737	GLYCOPROTEIN DEGRADATION
738	HYALURONOGLUCOSAMINIDASE
739	EXTRACELLULAR MATRIX MAINTENANCE
740	SERINE CARBOXYPEPTIDASE
741	ION CHANNEL
742	ION TRANSPORTER
743	SENSORY PERCEPTION
744	PAIN SENSATION
745	THERMORECEPTION AND RESPONSE
746	CYTOSOLIC RIBOSOME
747	L-LACTATE DEHYDROGENASE
748	HETEROTRIMERIC G-PROTEIN GTPASE, GAMMA SUBUNIT
749	RAB SMALL MONOMERIC GTPASE
750	RNA POLYMERASE I TRANSCRIPTION FACTOR
751	C-5 STEROL DESATURASE
752	CATABOLISM
753	CARBOXYLESTERASE
754	ADDICTION
755	VITAMIN B12 TRANSPORT
756	PHYSIOLOGICAL PROCESSES
757	VITAMIN BIOSYNTHESIS
758	CALCIUM ION HOMEOSTASIS
759	CALCIDIOL 1-MONOXYGENASE
760	OOGENESIS
761	CYSTEINE-TYPE PEPTIDASE
	G-PROTEIN COUPLED RECEPTOR PROTEIN SIGNALING
762	PATHWAY
763	PROTEIN TYROSINE/THREONINE PHOSPHATASE
764	DYNACTIN COMPLEX
765	INTEGRIN LIGAND
766	INTEGRIN RECEPTOR SIGNAL SIGNALLING PATHWAY
	NEGATIVE REGULATION OF HOMEOTIC GENE (POLYCOMB GROUP)
767	

768	FOCAL ADHESION
769	SPECTRIN
770	CELL DEATH
771	FERRITIN
772	IRON BINDING
773	PEPTIDYLPROLYL ISOMERASE
774	MICROTUBULE CYTOSKELETON
	GAMMA-AMINOBUTYRIC ACID-INHIBITED CHLORIDE
775	CHANNEL
776	GLUTATHIONE SYNTHASE
777	AMINO ACID METABOLISM
778	TRANSMEMBRANE RECEPTOR
779	EXCRETION
780	TRANSCRIPTION REGULATION, FROM POL II PROMOTER
781	CASPASE ACTIVATION
782	STAT PROTEIN DIMERIZATION
783	NIK-I-KAPPAB/NF-KAPPAB CASCADE
784	RESPONSE TO PEST/PATHOGEN/PARASITE
785	STAT PROTEIN NUCLEAR TRANSLOCATION
786	TYROSINE PHOSPHORYLATION OF STAT PROTEIN
787	METABOTROPIC GLUTAMATE RECEPTOR
	METABOTROPHIC GLUTAMATE RECEPTOR, PHOSPHOLIPASE
788	C ACTIVATING PATHWAY
789	HUMORAL DEFENSE MECHANISM
790	INTERCELLULAR JUNCTION
791	PEROXISOME-ASSEMBLY ATPASE
792	PHOSPHATE CARRIER
793	MITOCHONDRIAL INNER MEMBRANE
794	PROTEIN KINASE C BINDING PROTEIN
795	MITOTIC S-SPECIFIC TRANSCRIPTION
796	INDUCTION OF APOPTOSIS BY DNA DAMAGE
797	RECOMBINATION
798	AXON GUIDANCE RECEPTOR
799	COPII VESICLE
800	SNRNA TRANSCRIPTION
801	PROTEIN DEGRADATION TAGGING
802	ELECTRON TRANSPORT
803	AEROBIC RESPIRATION
804	RESPIRATORY CHAIN COMPLEX
805	CYTOCHROME C OXIDASE BIOGENESIS
806	TFIID COMPLEX
807	CYANATE CATABOLISM
808	THIOSULFATE SULFURTRANSFERASE
809	STRIATED MUSCLE CONTRACTION REGULATION

810	ENHANCER BINDING
811	MICROTUBULE CYTOSKELETON ORGANIZATION AND BIOGENESIS
812	RHO GUANYL-NUCLEOTIDE EXCHANGE FACTOR
813	LACTOSE BIOSYNTHESIS
814	PROGENY NUTRITION
815	VASCULAR ENDOTHELIAL GROWTH FACTOR RECEPTOR
816	PHOSPHOLIPASE INHIBITOR
817	APOLipoprotein
818	LOW-DENSITY LIPOPROTEIN
819	INTRACELLULAR IRON STORAGE
820	OTHER
821	SCAVENGER RECEPTOR
822	RNA CATABOLISM
823	POLY(A) BINDING
824	RNA TURNOVER
825	LDL RECEPTOR
826	LIPOPROTEIN BINDING
827	O-LINKED GLYCOSYLATION
828	LOW-DENSITY LIPOPROTEIN RECEPTOR
829	PURINE BASE METABOLISM
830	RIBOSE-PHOSPHATE PYROPHOSPHOKINASE
831	CARBOXYPEPTIDASE
832	INTRACELLULAR TRANSPORTER
833	ESTERASE, UNKNOWN SUBSTRATE
834	EARLY ENDOSOME
835	VESICLE FUSION
836	GLUCOSE TRANSPORT
837	GLUCOSE TRANSPORTER
838	INTERNALIZATION RECEPTOR
839	RECEPTOR MEDIATED ENDOCYTOSIS
840	CARBOXYPEPTIDASE D
841	DEOXYRIBONUCLEASE
842	TGF β RECEPTOR COMPLEX ASSEMBLY
843	INTERFERON-GAMMA RECEPTOR
844	RESISTANCE TO PATHOGENIC BACTERIA
845	SPHINGOMYELIN METABOLISM
846	SPHINGOMYELIN PHOSPHODIESTERASE
847	POST-REPLICATION REPAIR
848	DIGESTION
849	PEPTIDE HORMONE
850	NUTRITIONAL RESPONSE PATHWAY
851	INDUCTION OF APOPTOSIS BY HORMONES
852	ALPHA-MANNOSIDASE

853	PROTEIN DEGLYCOSYLATION
854	PHOSPHORIBOSYLAMINOIMIDAZOLE-SUCCINOCARBOXAMIDE SYNTHASE
855	BETA-N-ACETYLHEXOSAMINIDASE
856	ALKYL HYDROPEROXIDE REDUCTASE
857	ACTIN MONOMER BINDING
	TRANSMEMBRANE RECEPTOR PROTEIN TYROSINE PHOSPHATASE
858	PROTEOGLYCAN
860	ENZYME
861	CERAMIDE METABOLISM
862	LATE ENDOSOME
863	HYALURONIC ACID BINDING
864	COPPER BINDING
865	COPPER HOMEOSTASIS
866	COPPER ION TRANSPORT
867	mRNA CAP BINDING
868	PROTEIN KINASE C INHIBITOR
869	CAMP-DEPENDENT PROTEIN KINASE
870	ADENINE TRANSPORTER
871	MITOCHONDRIAL GENOME MAINTENANCE
872	HEPARIN BINDING
873	PROCOLLAGEN-LYSINE 5-DIOXYGENASE
874	MALE GONAD DEVELOPMENT
875	MALE SPECIFIC DEVELOPMENT
876	NUCLEOSIDE METABOLISM
877	NUCLEOSIDE-DIPHOSPHATE KINASE
878	GLYCOSAMINOGLYCAN CATABOLISM
879	COPPER-EXPORTING ATPASE
880	GOLGI TRANS-FACE
881	GOLGI-PLASMA MEMBRANE TRANSPORT VESICLE
882	TRANSCRIPTION FACTOR TFIIH
883	HEMOCYTE DEVELOPMENT
	TRANSMEMBRANE RECEPTOR PROTEIN TYROSINE KINASE
884	ADAPTOR PROTEIN
885	DNA REPLICATION INITIATION
886	ALDEHYDE REDUCTASE
887	FAT BODY DEVELOPMENT
888	TRIACYLGLYCEROL METABOLISM
889	DIACYLGLYCEROL O-ACYLTRANSFERASE
890	CASEIN KINASE I
891	ROUGH ENDOPLASMIC RETICULUM
892	DOUBLE-STRANDED RNA BINDING
893	UV RESPONSE

894	TRANSCRIPTION INITIATION
895	DNA-NONHOMOLOGOUS END-JOINING
896	SODIUM TRANSPORT
897	SODIUM:PHOSPHATE SYMPORTER
898	CIRCADIAN RHYTHM
899	LOCOMOTORY BEHAVIOR
900	NEUROPEPTIDE Y RECEPTOR
901	CALCIUM CHANNEL REGULATOR
	G-PROTEIN SIGNALLING, ADENYLYL CYCLASE INHIBITING
902	PATHWAY
903	FEEDING
904	LOCOMOTION
905	RIBONUCLEASE P
906	PROTEIN-NUCLEUS IMPORT, DOCKING
907	EXIT FROM MITOSIS
908	SEPTIN ASSEMBLY AND SEPTUM FORMATION
909	RESPONSE TO INJURY
910	DELTA DNA POLYMERASE
911	INTERMEDIATE FILAMENT
912	CONTROL OF MITOSIS
913	SINGLE-STRANDED RNA BINDING
914	FORMATE-TETRAHYDROFOLATE LIGASE
915	CALCIUM ION TRANSPORT
916	N-ACETYLTRANSFERASE
917	INTERNAL PROTEIN ACETYLATION
918	MITOTIC SPINDLE ASSEMBLY
919	RAN SMALL MONOMERIC GTPASE
920	UBIQUITINYL HYDROLASE 1
921	KDEL RECEPTOR
922	STEROID BIOSYNTHESIS
923	CELL-SUBSTRATE JUNCTION ASSEMBLY
924	MILK PROTEIN
925	CITRATE LYASE
926	ATP CATABOLISM
927	CITRATE METABOLISM
928	COENZYME A METABOLISM
929	ATP-CITRATE (PRO-S)-LYASE
930	PROTEIN KINASE C ACTIVATION
931	PROTEIN KINASE INHIBITION
932	INSULIN-LIKE GROWTH FACTOR RECEPTOR
933	HEME BIOSYNTHESIS
934	PORPHOBILINOGEN SYNTHASE
935	PLASMA PROTEIN
936	THYROID HORMONE TRANSPORTER

937	CONSERVED ATPASE DOMAIN
938	UBIQUITIN ISOPEPTIDASE T
939	LYSOSOMAL CYSTEINE-TYPE ENDOPEPTIDASE
940	DNA LIGASE
941	GLYCOLYSIS
942	STRIATED MUSCLE CONTRACTION
943	FRUCTOSE-BISPHOSPHATE ALDOLASE
944	VOLTAGE-GATED POTASSIUM CHANNEL
945	INWARD RECTIFIER POTASSIUM CHANNEL
946	MACROPHAGE ELASTASE
947	CONNECTIVE TISSUE DEVELOPMENT AND MAINTENANCE
948	SERINE-TYPE PEPTIDASE
949	7S RNA BINDING
950	3'-5' EXORIBONUCLEASE
951	PROTEIN C (ACTIVATED)
952	IRON-SULFUR ELECTRON TRANSFER CARRIER
953	MHC-INTERACTING PROTEIN
954	CELL CYCLE CHECKPOINT
955	MYOBLAST DETERMINATION
956	IMPORTIN ALPHA EXPORT RECEPTOR
957	mRNA-NUCLEUS EXPORT
958	NUCLEOSOME ASSEMBLY CHAPERONE
959	DNA REPLICATION DEPENDENT NUCLEOSOME ASSEMBLY
960	MITOTIC CHROMOSOME
961	ANTIBACTERIAL RESPONSE PROTEIN
962	GLUTATHIONE PEROXIDASE
963	mRNA CATABOLISM
964	STEROID METABOLISM
965	C-4 METHYL STEROL OXIDASE
966	PEROXISOME TARGETING SIGNAL-2 RECEPTOR
967	ACTIN CROSS-LINKING
968	GLYCOSAMINOGLYCAN BIOSYNTHESIS
969	GRAM-NEGATIVE BACTERIAL BINDING
970	V-SNARE
971	ER-GOLGI TRANSPORT VESICLE
972	ACETYL-COA CARBOXYLASE
973	METHYLTRANSFERASE
974	PROTEIN METHYLATION
975	CYTOSKELETAL ANCHORING
976	MEIOTIC PROPHASE II
977	UDP-GLUCOSE METABOLISM
978	UDP-GLUCOSE 6-DEHYDROGENASE
979	UDP-GLUCURONATE BIOSYNTHESIS
980	URIDINE KINASE

981	PYRIMIDINE RIBONUCLEOTIDE BIOSYNTHESIS
982	GDP-MANNOSE 4,6-DEHYDRATASE
983	HEXOKINASE
984	GLUCOSE METABOLISM
985	AMILORIDE-SENSITIVE SODIUM CHANNEL
986	OSMOREGULATION AND EXCRETION
987	2,4-DIEENOYL-COA REDUCTASE (NADPH)
988	KATANIN
989	MICROTUBULE-SEVERING ATPASE
990	MICROTUBULE DEPOLYMERIZATION
991	3-OXOACYL-[ACYL-CARRIER PROTEIN] REDUCTASE
992	MEMBRANE ASSOCIATED ACTIN BINDING
993	ADENINE PHOSPHORIBOSYLTRANSFERASE
994	tRNA BINDING
995	tRNA PROCESSING
996	ALANYL-tRNA BIOSYNTHESIS
997	ALCOHOL METABOLISM
998	ALDEHYDE DEHYDROGENASE (NAD+)
999	ALDEHYDE DEHYDROGENASE (NAD(P)+)
1000	ASPARAGINE-tRNA LIGASE
1001	CALCIUM STORAGE
1002	ENDOPLASMIC RETICULUM LUMEN
1003	HEMOGLOBIN
1004	FATTY ACID (OMEGA-1)-HYDROXYLASE
1005	INORGANIC DIPHOSPHATASE
1006	ISOCITRATE METABOLISM
1007	ISOCITRATE DEHYDROGENASE (NADP+)
1008	DIHYDROPRIMIDINASE
1009	GLYCOGEN CATABOLISM
1010	COCHAPERONIN
1011	ECTODERM DEVELOPMENT
1012	LECTIN
1013	METHIONINE ADENOSYLTRANSFERASE
1014	ER RETENTION
1015	PROTEIN DISULFIDE ISOMERASE
1016	PROTEIN TRANSPORTER
1017	THIOREDOXIN PEROXIDASE
1018	UBIQUITIN
1019	POLY-UBIQUITIN
1020	CYTOKINE
1021	PROTEIN PROLINE HYDROXYLATION
	PROCOLLAGEN-PROLINE,2-OXOGLUTARATE-4-
1022	DIOXYGENASE
1023	DNA MODIFICATION

1024	PURINE-NUCLEOSIDE PHOSPHORYLASE
1025	PYRUVATE KINASE
1026	ARGINYL-TRNA BIOSYNTHESIS
1027	THIOREDOXIN
1028	PLASMA GLYCOPROTEIN
1029	POLYUBIQUITYLATION
1030	DNA REPAIR REGULATION
1031	CHLORIDE CHANNEL
1032	HEART DEVELOPMENT
1033	POTASSIUM CHANNEL REGULATOR
1034	LIGAND-GATED ION CHANNEL
1035	CYSTINE TRANSPORTER
1036	AMINO ACID TRANSPORTER
	DOLICHYL-DIPHOSPHOOLIGOSACCHARIDE-PROTEIN
1037	GLYCOSYLTRANSFERASE
1038	GALACTOSYLTRANSFERASE
1039	DEATH RECEPTOR INTERACTING PROTEIN
1040	RESPONSE TO WOUNDING
1041	GLUTAMINE-TRNA LIGASE
1042	GLUTAMINYL-TRNA BIOSYNTHESIS
1043	POTASSIUM CHANNEL
1044	3-PHOSPHOINOSITIDE-DEPENDENT PROTEIN KINASE
1045	DNA DEPENDENT ADENOSINETRIPHOSPHATASE
1046	MICROVILLI
1047	ADENYLYL CYCLASE ACTIVATION
1048	ESTABLISHMENT OF CELL POLARITY
1049	CELL POLARITY
1050	PROTEASOME ATPASE
1051	CITRATE (SI)-SYNTHASE
1052	ISOVALERYL-COA DEHYDROGENASE
1053	COLD RESPONSE
1054	PERINUCLEAR SPACE
1055	MYOCYTE FUSION
1056	VOLTAGE-GATED CALCIUM CHANNEL
1057	LOW VOLTAGE-GATED CALCIUM CHANNEL
1058	INTERLEUKIN-5 RECEPTOR
1059	BETA-TUBULIN FOLDING
1060	TRANSLATION RELEASE FACTOR
1061	TRANSLATION TERMINATION FACTOR
1062	TRANSLATIONAL REGULATION, TERMINATION
1063	TRIOSEPHOSPHATE ISOMERASE
1064	NON-SELECTIVE VESICLE ASSEMBLY
1065	ARF SMALL MONOMERIC GTPASE
1066	FERTILIZATION

1067	ACETYL-COA C-ACETYLTRANSFERASE
1068	FATTY ACID BETA-OXIDATION
1069	LONG-CHAIN ACYL-COA DEHYDROGENASE
	ENERGY DERIVATION BY OXIDATION OF ORGANIC
1070	COMPOUNDS
1071	CARBONATE DEHYDRATASE
1072	MEMBRANE DIPEPTIDASE
1073	RETINOID BINDING
1074	ARGININE CATABOLISM
1075	GUANYL-NUCLEOTIDE EXCHANGE FACTOR
1076	FUMARATE HYDRATASE
1077	FUMARATE METABOLISM
1078	HEMOSTASIS
1079	GLUCOSE 6-PHOSPHATE UTILIZATION
1080	GLUCOSE-6-PHOSPHATE 1-DEHYDROGENASE
1081	CYSTEINE METABOLISM
1082	GLUTAMATE METABOLISM
1083	GLUTATHIONE BIOSYNTHESIS
1084	GLUTAMATE-CYSTEINE LIGASE
1085	AT DNA BINDING
1086	THIOREDOXIN REDUCTASE (NADPH)
1087	CHROMATIN ASSEMBLY/DISASSEMBLY
1088	KILLER ACTIVITY
1089	RESPIRATION
1090	PENTOSE-PHOSPHATE SHUNT, OXIDATIVE BRANCH
1091	HSP70/HSP90 ORGANIZING PROTEIN
1092	DNA FRAGMENTATION
1093	PHOSPHATIDYLETHANOLAMINE BINDING
1094	PROTEIN TYROSINE PHOSPHATASE ACTIVATOR
1095	PYRROLINE 5-CARBOXYLATE REDUCTASE
1096	SMOOTH ENDOPLASMIC RETICULUM
1097	SMALL UBIQUITIN-RELATED PROTEIN 1 CONJUGATION
1098	BRUSH BORDER
1099	CREATINE KINASE
1100	PENTOSE-PHOSPHATE SHUNT
1101	ALDEHYDE DEHYDROGENASE
1102	PERIPHERAL NERVOUS SYSTEM DEVELOPMENT
1103	ENOYL-COA HYDRATASE
1104	ACETYL-COA C-ACYLTRANSFERASE
1105	3-HYDROXYACYL-COA DEHYDROGENASE
1106	NITRIC OXIDE SYNTHASE
1107	CLATHRIN ADAPTOR
1108	EXTRACELLULAR MATRIX GLYCOPROTEIN
1109	HIGH DENSITY LIPOPROTEIN BINDING

1110	CALCIUM CHANNEL
1111	GLUTATHIONE REDUCTASE (NADPH)
1112	HOMOPHILIC CELL ADHESION
1113	CALCIUM-INDEPENDENT CELL-CELL MATRIX ADHESION
1114	EMBRYONIC POLARITY
1115	MYO-INOSITOL:SODIUM SYMPORTER
1116	BEHAVIOR
1117	CANNABINOID RECEPTOR
	G-PROTEIN SIGNALLING, LINKED TO CYCLIC NUCLEOTIDE
1118	SECOND MESSENGER
1119	TIGHT JUNCTION
	MEMBRANE-ASSOCIATED PROTEIN WITH GUANYLATE
1120	KINASE ACTIVITY
1121	PERICENTRIOLAR MATERIAL
1122	FOCAL ADHESION KINASE
1123	SIGNAL COMPLEX FORMATION
1124	UREA CYCLE
1125	ARGININOSUCCINATE LYASE
1126	NUCLEOTIDE-SUGAR METABOLISM
1127	MEDIATOR COMPLEX
1128	FATTY-ACYL-COA SYNTHASE
1129	TRANSCRIPTION TERMINATION
1130	TRANSCRIPTION ELONGATION FACTOR COMPLEX
1131	POL II TRANSCRIPTION TERMINATION FACTOR
1132	CASPASE-3
1133	POLY(ADP-RIBOSE) GLYCOHYDROLASE
1134	VOLTAGE-SENSITIVE CALCIUM CHANNEL
1135	tRNA GUANYLYLTRANSFERASE
1136	GLYCOSAMINOGLYCAN BINDING
1137	PROTEIN SERINE/THREONINE PHOSPHATASE
1138	RHO GTPASE ACTIVATOR
1139	CYTOSKELETON ORGANIZATION AND BIOGENESIS
1140	DNA LIGATION
1141	ALDEHYDE METABOLISM
1142	ALDO-KETO REDUCTASE
1143	CALCIUM-DEPENDENT PHOSPHOLIPID BINDING
	DIPHOSPHOINOSITOL POLYPHOSPHATE
1144	PHOSPHOHYDROLASE
1145	HETEROTRIMERIC G PROTEIN
1146	T CELL RECEPTOR
1147	POLY(U) BINDING
1148	ACETYL-COA METABOLISM
1149	CYTOSKELETAL ADAPTOR
1150	CAM-DEPENDENT CYCLIC-NUCLEOTIDE

	PHOSPHODIESTERASE
1151	CYCLIN-DEPENDENT PROTEIN KINASE INHIBITOR
1152	SERINE BIOSYNTHESIS
1153	PHOSPHOGLYCERATE DEHYDROGENASE
1154	TRYPSIN
1155	MITOTIC S PHASE
	G-PROTEIN-COUPLED RECEPTOR PHOSPHORYLATING
1156	PROTEIN KINASE
1157	OXIDATIVE PHOSPHORYLATION
1158	SMOOTH MUSCLE CONTRACTION REGULATION
1159	METHYLMALONYL-COA MUTASE
1160	CHEMOSENSATION AND RESPONSE
1161	CARBAMOYL-PHOSPHATE SYNTHASE (AMMONIA)
1162	NUCLEAR RNA-NUCLEUS EXPORT
1163	EMBRYO IMPLANTATION
1164	SMALL RIBOSOMAL SUBUNIT
1165	ENDOPLASMIC RETICULUM RECEPTOR
1166	DNA BENDING
1167	INTRACELLULAR
1168	PROTEIN SYNTHESIS INITIATION
1169	ION TRANSPORT
1170	GUANYL-NUCLEOTIDE RELEASING FACTOR
1171	SPliceosome ASSEMBLY
1172	ADENYLYL CYCLASE
	CALCIUM/CALMODULIN-RESPONSIVE ADENYLYL
1173	CYCLASE
1174	NASCENT POLYPEPTIDE ASSOCIATION
1175	NASCENT POLYPEPTIDE-ASSOCIATED COMPLEX
1176	NONSENSE-MEDIATED mRNA DECAY
1177	ACYL-COA METABOLISM
1178	DEGRADATION OF CYCLIN
1179	CYCLIN SELECTIVE UBIQUITIN CONJUGATING ENZYME
1180	STEROID BINDING
1181	GLYCOSPHINGOLIPID METABOLISM
1182	MULTIDRUG TRANSPORTER
1183	ORGANIC ANION TRANSPORTER
1184	ASIALOGLYCOPEPTIDE RECEPTOR
1185	NUCLEAR INNER MEMBRANE
1186	RIBONUCLEASE INHIBITOR
1187	CALCIUM/CALMODULIN-DEPENDENT PROTEIN KINASE
1188	NEUTRAL AMINO ACID TRANSPORT
1189	NEUTRAL AMINO ACID TRANSPORTER
1190	TRANSPORT
1191	CALPAIN

1192	PROTEIN PHOSPHATASE TYPE 2A CATALYST
1193	MEMORY
1194	MRNA SPlice SITE SELECTION
1195	DNA TOPOISOMERASE I
1196	DRUG RESISTANCE
1197	KINESIN
1198	MICROTUBULE MOTOR
1199	NADPH-FERRIHEMOPROTEIN REDUCTASE
1200	MRNA BINDING, 3' UTR
1201	COATOMER
	EUKARYOTIC TRANSLATION INITIATION FACTOR 2ALPHA
1202	KINASE
1203	HEAT RESPONSE
	RECEPTOR SIGNALLING PROTEIN SERINE/THREONINE
1204	KINASE
1205	NUCLEASE
1206	RNA MODIFICATION
1207	GOLGI VESICLE
1208	TYROSINE-TRNA LIGASE
1209	TYROSYL-TRNA BIOSYNTHESIS
1210	INTERLEUKIN-8 RECEPTOR LIGAND
1211	PURINE SALVAGE
1212	HYPOXANTHINE PHOSPHORIBOSYLTRANSFERASE
1213	HEME OXYGENASE (DECYCLIZING)
1214	TRNA MODIFICATION
1215	HISTONE MRNA METABOLISM
1216	MAPKKK CASCADE
1217	FGF RECEPTOR SIGNALLING PATHWAY
1218	FIBROBLAST GROWTH FACTOR RECEPTOR
1219	RAS GUANYL-NUCLEOTIDE EXCHANGE FACTOR
1220	PHOSPHOGLYCERATE MUTASE
1221	GLUTATHIONE TRANSFERASE
1222	HEAVY METAL RESISTANCE
1223	HEAVY METAL RESPONSE
1224	HEAVY METAL ION TRANSPORT
1225	COPPER, ZINC SUPEROXIDE DISMUTASE
1226	CYTOPLASMIC VESICLE
1227	CELL ADHESION INHIBITION
1228	RHO GDP-DISSOCIATION INHIBITOR
1229	CELL FATE SPECIFICATION
1230	CILIUM
1231	MORPHOGENESIS
1232	PHOSPHATIDYLINOSITOL TRANSPORTER
1233	COCHAPERONE

1234	POST-CHAPERONINE TUBULIN FOLDING PATHWAY
1235	PROTEINASE INHIBITOR
1236	HEAVY METAL ION TRANSPORTER
1237	LACTASE
1238	PHOSPHOFRUCTOKINASE
1239	GLYCOLYSIS REGULATION
1240	SEPTATE JUNCTION
1241	CELL-CELL ADHERENS JUNCTION
1242	INTERCELLULAR JUNCTION ASSEMBLY
1243	LACTOYLGLUTATHIONE LYASE
1244	HYDROXYMETHYLBILANE SYNTHASE
1245	DUTP PYROPHOSPHATASE
1246	PROTEIN PHOSPHATASE TYPE 2C
1247	INTERFERON-ALPHA/BETA RECEPTOR
1248	GLYCINE METABOLISM
1249	BILE ACID METABOLISM
1250	ARYLSULFATASE
1251	LYSOSOMAL TRANSPORT
1252	HYDROGEN-TRANSLOCATING V-TYPE ATPASE
1253	HEME TRANSPORTER
1254	GLYCOGEN PHOSPHORYLASE
1255	CREATINE TRANSPORTER
1256	NEUROTRANSMITTER UPTAKE
1257	CREATINE:SODIUM SYMPORter
1258	EICOSANOID METABOLISM
1259	CALCIUM-DEPENDENT CYTOSOLIC PHOSPHOLIPASE A2
1260	ENDORIBONUCLEASE
1261	ALDEHYDE OXIDASE
1262	XANTHINE DEHYDROGENASE
1263	OXYGEN AND RADICAL METABOLISM
1264	BLOOD GROUP ANTIGEN
1265	OXYGEN TRANSPORTER
1266	OXYGEN TRANSPORT
1267	NITRILASE
1268	RENIN
1269	DNA DAMAGE INDUCED PROTEIN PHOSPHORYLATION
1270	BLASTODERM SEGMENTATION
1271	MEMBRANE PROTEIN ECTODOMAIN PROTEOLYSIS
1272	SPINDLE MICROTUBULE
1273	ANTEROGRADE AXON CARGO TRANSPORT
1274	ORGANELLE ORGANIZATION AND BIOGENESIS
1275	GLYCIPAN
1276	EXO-ALPHA-SIALIDASE
1277	MANNOSYLTRANSFERASE

1278	LIPOPOLYSACCHARIDE BIOSYNTHESIS
1279	METABOLISM
1280	GPI-ANCHOR TRANSAMIDASE
1281	TROPOMYOSIN
1282	MUSCLE CONTRACTION REGULATION
1283	SYNAPTIC VESICLE
1284	NEUROTRANSMITTER RELEASE
1285	GOLGI STACK
1286	GLUTAREDOXIN
1287	ACID PHOSPHATASE
1288	DOPACHROME DELTA-ISOMERASE
1289	SUBSTRATE-BOUND CELL MIGRATION, CELL EXTENSION
1290	INTRACELLULAR COPPER DELIVERY
1291	CATHEPSIN D
1292	LEUKOTRIENE METABOLISM
1293	SPERMIDINE SYNTHASE
1294	POLYAMINE METABOLISM
1295	METHIONINE METABOLISM
1296	PROTEIN PHOSPHATASE INHIBITOR
1297	DEATH RECEPTOR LIGAND
1298	APOPTOTIC MITOCHONDRIAL CHANGES
1299	INDUCTION OF APOPTOSIS VIA DEATH DOMAIN RECEPTORS
1300	MOLECULAR_FUNCTION
1301	GLYCEROPHOSPHOLIPID METABOLISM
1302	1-PHOSPHATIDYLINOSITOL-4-PHOSPHATE KINASE
1303	MALATE DEHYDROGENASE
1304	CALCIUM-DEPENDENT CELL ADHESION
1305	THIOPURINE S-METHYLTRANSFERASE
1306	RHO SMALL MONOMERIC GTPASE
1307	ADP REDUCTION
1308	NUCLEOTIDE METABOLISM
1309	RIBONUCLEOSIDE DIPHOSPHATE CATABOLISM
1310	RIBONUCLEASE
1311	FATTY ACID BINDING
1312	MOLECULAR_FUNCTION UNKNOWN
1313	GERM CELL DEVELOPMENT
1314	REPRESSION OF SURVIVAL GENE PRODUCTS
1315	DIADENOSINE POLYPHOSPHATE CATABOLISM
1316	EXONUCLEASE
1317	UV PROTECTION
	DOUBLE-STRANDED DNA SPECIFIC
1318	EXODEOXYRIBONUCLEASE
1319	EGF RECEPTOR MODULATION
1320	AXONEMAL MOTOR

1321	AXONEMAL DYNEIN
1322	PROTEIN PHOSPHATASE
1323	INOSITOL/PHOSPHATIDYLINOSITOL PHOSPHATASE
1324	OSSIFICATION
1325	GLUCOSAMINE CATABOLISM
1326	GLUCOSAMINE-6-PHOSPHATE ISOMERASE
1327	MITOTIC METAPHASE/ANAPHASE TRANSITION
1328	SELENIUM BINDING
1329	G/T-MISMATCH-SPECIFIC THYMINE-DNA GLYCOSYLASE
1330	NON-SELECTIVE VESICLE DOCKING
1331	INTRA GOLGI TRANSPORT
1332	INTER-GOLGI TRANSPORT VESICLE
1333	RAS SMALL MONOMERIC GTPASE
1334	PHOSPHOMEVALONATE KINASE
1335	PROTEASOME ACTIVATOR
1336	THYMIDYLATE KINASE
1337	PROSTAGLANDIN METABOLISM
1338	SINGLE-STRAND BREAK REPAIR
1339	PHOSPHATIDYLCHOLINE TRANSPORTER
1340	NEUROTRANSMITTER SYNTHESIS AND STORAGE
1341	TRANSALDOLASE
1342	SYNAPTONEMAL COMPLEX
1343	DIHYDROLIPOAMIDE DEHYDROGENASE
1344	CATABOLIC CARBOHYDRATE METABOLISM
1345	DEATH RECEPTOR ASSOCIATED FACTOR
1346	HYDROGEN TRANSPORTER
1347	PHENYLALANINE METABOLISM
1348	TETRAHYDROBIOPTERIN BIOSYNTHESIS
1349	4A-HYDROXYTETRAHYDROBIOPTERIN DEHYDRATASE
1350	GALACTOKINASE
1351	GALACTOSE METABOLISM
1352	BIS(5'-NUCLEOSYL)-TETRAPHOSPHATASE (SYMMETRICAL)
1353	IONIC INSULATION OF NEURONS BY GLIAL CELLS TYPE 1 SERINE/THREONINE SPECIFIC PROTEIN
1354	PHOSPHATASE INHIBITOR
1355	BIOLOGICAL PROCESS UNKNOWN
1356	CATHEPSIN H
1357	CASPASE-ACTIVATED DEOXYRIBONUCLEASE
1358	ACYLPHOSPHATASE
1359	ACYL-COA BINDING
1360	PROLYL OLIGOPEPTIDASE
1361	GROWTH FACTOR
1362	PHOSPHATIDYLINOSITOL-BISPHEROSPATASE
1363	ION CHANNEL INHIBITOR

1364	DIHYDROBIOPTERIN REDUCTION
1365	DIHYDROPTERIDINE REDUCTASE
1366	DIAZEPAM-BINDING INHIBITOR
1367	GALACTOSE BINDING LECTIN
1368	ORNITHINE METABOLISM
1369	ORNITHINE-OXO-ACID AMINOTRANSFERASE
1370	CATHEPSIN B
1371	BILE ACID TRANSPORTER
1372	CTP SYNTHASE
1373	SORBITOL METABOLISM
1374	UBIQUITIN-LIKE ACTIVATING ENZYME
1375	DIHYDROLIPOAMIDE S-ACETYLTRANSFERASE
1376	PHOSPHORYLASE
1377	GLUTAMATE CATABOLISM
1378	FRUCTOSE 2,6-BISPHOSPHATE METABOLISM
1379	FRUCTOSE-2,6-BISPHOSPHATE 2-PHOSPHATASE
1380	3-BETA-HYDROXY-DELTA(5)-STEROID DEHYDROGENASE
1381	ALPHA DNA POLYMERASE:PRIMASE COMPLEX
1382	ACONITATE HYDRATASE
1383	MITOCHONDRIAL LARGE RIBOSOMAL-SUBUNIT
1384	mRNA EDITING
1385	CYCLOSPORIN A BINDING
1386	MEVALONATE TRANSPORT
1387	MEVALONATE TRANSPORTER
1388	MONOCARBOXYLIC ACID TRANSPORT
1389	MONOCARBOXYLIC ACID TRANSPORTER
1390	P-ELEMENT BINDING
1391	AMINOACYLASE
1392	ASPARTATE-TRNA LIGASE
1393	ASPARTYL-TRNA BIOSYNTHESIS
1394	POLYPEPTIDE N-ACETYLGLACTOSAMINYLTRANSFERASE
1395	COLLAGEN
1396	HYDROGEN/POTASSIUM-EXCHANGING ATPASE
1397	SARCOGLYCAN COMPLEX
1398	FUCOSYLTRANSFERASE
1399	AMINOPEPTIDASE
1400	UDP-GLUCOSE 4-EPIMERASE
1401	TRANSPORTIN
1402	METHIONYL AMINOPEPTIDASE
1403	DIPEPTIDYL-PEPTIDASE
1404	PITRILYSIN
1405	2',3'-CYCLIC NUCLEOTIDE 3'-PHOSPHODIESTERASE
1406	NUCLEAR OUTER MEMBRANE
1407	IMP DEHYDROGENASE

1408	PEPTIDE METABOLISM
1409	SECRETORY VESICLE MEMBRANE
1410	GLYCINE-TRNA LIGASE
1411	HETEROTRIMERIC G-PROTEIN GTPASE, ALPHA SUBUNIT
1412	ARF GTPASE ACTIVATOR
1413	PEROXISOMAL MATRIX
1414	VERY LONG CHAIN FATTY ACID METABOLISM
1415	ALPHA-GLUCOSIDASE
1416	PHAGOCYTOSIS
1417	CREATINE BIOSYNTHESIS
1418	GLYCINE AMIDINOTRANSFERASE
1419	MAP KINASE
1420	CALCIUM-ACTIVATED POTASSIUM CHANNEL
1421	BLEOMYCIN HYDROLASE
1422	AMINOBUTYRATE CATABOLISM
1423	SUCCINATE-SEMALDEHYDE DEHYDROGENASE
1424	LYSOSOMAL MEMBRANE
1425	PHAGOSOME FORMATION
1426	DEBRANCHING ENZYME
1427	GLUTAMATE DEHYDROGENASE
1428	SPHINGOLIPID METABOLISM
1429	SERINE C-PALMITOYLTRANSFERASE
1430	ENDOSOME TO LYSOSOME TRANSPORT
1431	CELL GROWTH AND/OR MAINTENANCE
1432	NAD(P)(+)-TRANSHYDROGENASE (B-SPECIFIC)
1433	MAJOR HISTOCOMPATIBILITY PEPTIDE TRANSPORTER
1434	AMINOGLYCAN BIOSYNTHESIS
1435	ACETYLGLUCOSAMINYLTRANSFERASE
1436	UBIQUINOL-CYTOCHROME-C REDUCTASE
1437	ADENYLYLATE CYCLASE INHIBITION
1438	GLUCONEOGENESIS
1439	DICARBOXYLIC ACID TRANSPORT
1440	DICARBOXYLIC ACID TRANSPORTER
1441	CARRIER
1442	NON-SELECTIVE VESICLE FUSION
1443	MITOCHONDRIAL CITRATE TRANSPORT
	SODIUM:DICARBOXYLATE/TRICARBOXYLATE
1444	COTRANSPORTER
1445	SUBTILISIN
1446	PROPROTEIN CONVERTASE 2
1447	SERINE-TRNA LIGASE
1448	METALLOEXOPEPTIDASE
1449	HOLOCYTOCHROME C SYNTHASE
1450	AXON GUIDANCE

1451	PROTEIN-MEMBRANE TARGETING
1452	COMPLEMENT COMPONENT
1453	COMPLEMENT ACTIVATION
1454	OXOGLUTARATE DEHYDROGENASE (LIPOAMIDE)
1455	POTASSIUM:CHLORIDE SYMPORTER
1456	GLYCINE CATABOLISM
1457	GLYCINE DEHYDROGENASE (DECARBOXYLATING)
1458	OLIGOSACCHARIDE METABOLISM
1459	MANNOSYL-OLIGOSACCHARIDE 1,2-ALPHA-MANNOSIDASE
1460	N-GLYCAN PROCESSING
1461	UTP-GLUCOSE-1-PHOSPHATE URIDYLTRANSFERASE
1462	O-METHYLTRANSFERASE
1463	SOLUTE:CATION SYMPORTER
1464	CYTOCHROME C
1465	OXOGLUTARATE/MALATE ANTIPORTER
1466	3'-5' EXODEOXYRIBONUCLEASE
1467	CYTOCHROME B
1468	PYRUVATE METABOLISM
1469	MALATE DEHYDROGENASE (DECARBOXYLATING)
1470	PROTEIN KINASE C
1471	CASPASE ACTIVATION VIA CYTOCHROME C
1472	UDP-GLUCOSE:GLYCOPROTEIN GLUCOSYLTRANSFERASE
1473	ACTIVATION OF MAP/ERK KINASE KINASE
1474	CATALASE
1475	INSOLUBLE FRACTION
1476	MITOCHONDRIAL TRANSPORT
1477	GOLGI LUMEN
1478	ENDOCYTOTIC TRANSPORT VESICLE
1479	SODIUM DEPENDENT MULTIVITAMIN TRANSPORTER
1480	POLY-GLUTAMINE TRACT BINDING
1481	TRANSMEMBRANE RECEPTOR PROTEIN SERINE/THREONINE KINASE
1482	TRANSMEMBRANE RECEPTOR PROTEIN SERINE/THREONINE KINASE SIGNALLING PATHWAY
1483	DRUG TRANSPORTER
1484	STEROID DELTA-ISOMERASE
1485	PHOSPHORYLASE KINASE REGULATOR
1486	FERREDOXIN--NADP(+) REDUCTASE
1487	PROFILIN BINDING
1488	ATP-DEPENDENT PEPTIDASE
1489	EPOXIDE HYDROLASE
1490	LEUKOTRIENE-A4 HYDROLASE
1491	MANNOSE METABOLISM
1492	MANNOSE-6-PHOSPHATE ISOMERASE

1493	CELL WALL
1494	N-ACETYLGLUCOSAMINE METABOLISM
1495	N-ACETYLMANNOSAMINE METABOLISM
1496	CENTROMERE/KINETOCHEM COMPLEX MATURATION
1497	3-HYDROXYISOBUTYRYL-COA HYDROLASE
1498	PYRUVATE DEHYDROGENASE (LIPOAMIDE)
1499	COPROPORPHYRINOGEN OXIDASE
1500	EXOPEPTIDASE
1501	GAMMA-GLUTAMYL HYDROLASE
1502	DOUBLE-STRANDED RNA ADENOSINE DEAMINASE
1503	ETHANOL OXIDATION
1504	ETHANOL METABOLISM
1505	ALCOHOL DEHYDROGENASE
1506	ALCOHOL DEHYDROGENASE, ZINC-DEPENDENT
1507	ANTIMICROBIAL RESPONSE PROTEIN
1508	BRANCHED-CHAIN AMINO ACID AMINOTRANSFERASE
1509	BRANCHED CHAIN FAMILY AMINO ACID BIOSYNTHESIS
1510	SUCCINYL-COA METABOLISM
1511	3-OXOACID COA-TRANSFERASE
1512	PHOSPHATIDYLINOSITOL BIOSYNTHESIS
1513	PROTEIN PHOSPHATASE TYPE 1 REGULATOR
1514	SERINE METABOLISM
1515	PHOSPHOSERINE PHOSPHATASE
1516	MUTAGENESIS

A resulting entry for a peptide sequence based on these criteria will have the following format: SEQ ID NO of peptide entry | Numeric code corresponding to cell type and HLA type | SEQ ID NOs of source protein reference(s) | source protein symbol(s) | Numeric keys corresponding to biological classification(s). This ordering
5 corresponds to Criteria 1 | Criteria 2 | Criteria 3 | Criteria 4 | Criteria 5.

Example 2: Isolation and Characterization of Global Peptide Tags (GPT)

This example describes the use of enzymatic or chemical digestion strategies to reduce proteins of a complex mixture into peptides. These peptides are called Global
10 Peptide Tags (GPT). The peptides were separated and fractionated by multiple modes of chromatography and ultimately sequenced by liquid chromatography on-line with tandem mass spectrometry (LC/MS/MS).

Prior to digestion, all proteins of a sample were denatured using high concentrations of chemical denaturants (such as 6-8 M Urea or 6-8 M guanidine
15 hydrochloride), elevated temperature, or a combination of both chemical denaturants and elevated temperature. Additionally, reactive thiol groups were typically reduced by the action of dithiothreitol (DTT) or Tris[2-carboxyethylphosphine] hydrochloride (TCEP) at a molar concentration of 25 to 50 times greater than that of the total protein concentration, and alkylated with an alkylating agent (at a molar concentration of 25 to
20 50 times greater than that of the total protein concentration) such as iodoacetamide or iodoacetic acid. The reaction was typically carried out at room temperature and in the dark.

Two rounds of reduction and alkylation were usually used for each protein mixture to ensure complete reduction and alkylation of reactive thiol groups. Solutions
25 were typically made in a 50-100 mM ammonium bicarbonate solution at an approximate pH of 8.2. Subsequently, the reduced and alkylated protein mixture was concentrated to a volume of approximately 50-100 uL under vacuum in a Speedvac™ centrifugal concentrator (ThermoSavant Scientific) used at ambient temperature. This process also removes the majority of the excess DTT that is used to quench the final alkylation
30 reaction. Following concentration, the resultant solution was re-diluted with the ammonium bicarbonate solution and an enzyme was added at a weight to weight ratio of

1 part enzyme to 25-40 parts protein. Those protein mixtures that were chemically denatured were diluted with the ammonium bicarbonate solution to reduce the concentration of the chemical denaturant to less than 1 M. The enzyme was also dissolved in the ammonium bicarbonate solution. Enzymes that have been used to generate peptides from a complex protein mixture include trypsin and lysine endopeptidase. All enzymatic digestions were carried out overnight (typically 18 to 26 hours) at a temperature of 37°C. After protein digestion, enzymes were deactivated using 10% acetic acid solution, and peptides were separated from undigested protein and isolated by ultra filtration using either a 3 kDa or 5 kDa spin filtration device.

10 Alternatively, total protein digest are generated chemically using cyanogen bromide. Again, peptides were isolated from undigested proteins by ultra filtration. Peptide-rich solutions were separated by multiple modes of chromatography. The first mode was usually strong cation exchange (SCX) using a stationary phase such as polyethylaspartamide (from PolyLC Inc.) and an aqueous mobile phase that was modified with acetonitrile (5-15% v/v) and developed a salt gradient from 0 to 1 M salt to elute the adsorbed peptides. Each of the peptide-rich fractions that was isolated by SCX chromatography was further separated and fractionated using a C18 reversed phase microbore (1 mm id) column and mobile phases that were modified with trifluoroacetic acid and developed an acetonitrile gradient. Peptide-rich fractions that were isolated by reversed phase chromatography were subjected to on-line LC/MS/MS using a further dimension of reversed phase chromatography. Peptide sequence elucidation was by database searching raw MS/MS spectra against publicly available protein sequence databases.

25 Table 1 contains the sequences of a series of EPTs and GPTs identified as described in Examples 1 and 2. The conventions detailed in Example 1 are used to describe EPTs and GPTs, with the exception of "HLA source" which does not apply to GPTs.

TABLE 1

SEQ ID NO:1 2 SEQ ID NO:236, SEQ ID NO:237, SEQ ID NO:238, SEQ ID NO:239, SEQ ID NO:240, SEQ ID NO:241, SEQ ID NO:242, SEQ ID NO:243, SEQ ID NO:244, SEQ ID NO:245, SEQ ID NO:246 GRIN2A,
5 NMDAR2A 22,27,28,29,30,31,32,33,34
SEQ ID NO:2 2 SEQ ID NO:247, SEQ ID NO:248, SEQ ID NO:249 EFP, TRIM25, Z147, ZNF147 23,119,120
SEQ ID NO:3 2 SEQ ID NO:250, SEQ ID NO:251, SEQ ID NO:252, SEQ ID NO:253, SEQ ID NO:254, SEQ ID NO:255 GPM6, GPM6A, M6A 31,107
SEQ ID NO:4 2,10 SEQ ID NO:256, SEQ ID NO:257, SEQ ID NO:258, SEQ ID NO:259, SEQ ID NO:260, SEQ ID NO:261, SEQ ID NO:262, SEQ ID NO:263, SEQ ID NO:264 CD27L, CD27LG, CD70, KL-24 ANTIGEN, TNFSF7 31,56,57,89,120,179,387,388,389
10 SEQ ID NO:5 2,10 SEQ ID NO:265, SEQ ID NO:266, SEQ ID NO:267, SEQ ID NO:268, SEQ ID NO:269, SEQ ID NO:270, SEQ ID NO:271, SEQ ID NO:272, SEQ ID NO:273, SEQ ID NO:274, SEQ ID NO:275, SEQ ID NO:276, SEQ ID NO:277, SEQ ID NO:278, SEQ ID NO:279 CD132, IL2RG, IMD4, SCIDX, SCIDXL 31,34, 57,89,127,129,332,402,403,404
15 SEQ ID NO:6 2 SEQ ID NO:280, SEQ ID NO:281, SEQ ID NO:282, SEQ ID NO:283 BR, CD49B, ITGA2 17, 18,34,56,154,186,187,200,345,405,406,407,408,409,410
SEQ ID NO:7 2 SEQ ID NO:284, SEQ ID NO:285, SEQ ID NO:286, SEQ ID NO:287, SEQ ID NO:288, SEQ ID NO:289, SEQ ID NO:290, SEQ ID NO:291, SEQ ID NO:292, SEQ ID NO:293, SEQ ID NO:294 ERBB4, HER4 12,31,34,36,56,57,58,59,411
20 SEQ ID NO:8 2 SEQ ID NO:295, SEQ ID NO:296, SEQ ID NO:297, SEQ ID NO:298, SEQ ID NO:299 PA26 8,55,56,371,384,465
SEQ ID NO:9 9,24 SEQ ID NO:300, SEQ ID NO:301, SEQ ID NO:302, SEQ ID NO:303, SEQ ID NO:304, SEQ ID NO:305, SEQ ID NO:306, SEQ ID NO:307, SEQ ID NO:308, SEQ ID NO:309, SEQ ID NO:310, SEQ ID NO:311 CNND2, KIAK0002 62,136,137
25 SEQ ID NO:10 9 SEQ ID NO:312, SEQ ID NO:313, SEQ ID NO:314, SEQ ID NO:315, SEQ ID NO:316, SEQ ID NO:317, SEQ ID NO:318, SEQ ID NO:319, SEQ ID NO:320, SEQ ID NO:321, SEQ ID NO:322, SEQ ID NO:323, SEQ ID NO:324, SEQ ID NO:325, SEQ ID NO:326 ADPRT, ADPRT1, PADPRT1, PARP, PARP-1, PPOL 3, 7,8,10,12,13,18,53,60,179,335,336,337
30 SEQ ID NO:11 9,14 SEQ ID NO:327, SEQ ID NO:328, SEQ ID NO:329, SEQ ID NO:330, SEQ ID NO:331 ETFB 44,481,482
SEQ ID NO:12 9 SEQ ID NO:332, SEQ ID NO:333, SEQ ID NO:334, SEQ ID NO:335, SEQ ID NO:336, SEQ ID NO:337, SEQ ID NO:338, SEQ ID NO:339 DAGK5, DGKZ, HDGKZETA 8,12,89,98,362,487
35 SEQ ID NO:13 9,13,24,28 SEQ ID NO:340, SEQ ID NO:341, SEQ ID NO:342, SEQ ID NO:343, SEQ ID NO:352, SEQ ID NO:353, SEQ ID NO:354, SEQ ID NO:355, SEQ ID NO:356, SEQ ID NO:357, SEQ ID

NO : 358 , SEQ ID NO : 359 , SEQ ID NO : 360 | ACTR, AIB1, CAGH16, CTG26, NCOA3, P/CIP, RAC3, TNRC14, TNRC16 ,
 TRAM-1 | 8, 10, 17, 36, 89, 118, 151 |
 SEQ ID NO : 15 | 9, 13 | SEQ ID NO : 361 , SEQ ID NO : 362 , SEQ ID NO : 363 , SEQ ID NO : 364 , SEQ ID NO : 365 ,
 SEQ ID NO : 366 , SEQ ID NO : 367 , SEQ ID NO : 368 , SEQ ID NO : 369 , SEQ ID NO : 370 , SEQ ID NO : 371 , SEQ
 5 ID NO : 372 , SEQ ID NO : 373 | ASH2L, ASH2L2 | 7, 8, 10, 14, 53, 54 |
 SEQ ID NO : 16 | 9 | SEQ ID NO : 374 , SEQ ID NO : 375 , SEQ ID NO : 376 , SEQ ID NO : 377 , SEQ ID NO : 378 , SEQ
 ID NO : 379 , SEQ ID NO : 380 , SEQ ID NO : 381 , SEQ ID NO : 382 , SEQ ID NO : 383 , SEQ ID NO : 384 | CD53 ,
 MOX44 | 31, 56, 89, 129, 141, 154 |
 SEQ ID NO : 17 | 9 | SEQ ID NO : 385 , SEQ ID NO : 386 , SEQ ID NO : 387 , SEQ ID NO : 388 , SEQ ID NO : 389 |
 10 RGS14 | 17, 239, 242, 243, 441 |
 SEQ ID NO : 18 | 9, 13 | SEQ ID NO : 390 , SEQ ID NO : 391 , SEQ ID NO : 392 , SEQ ID NO : 393 , SEQ ID NO : 394 |
 DDX9, LKP, NDHII, RHA | 7, 8, 47, 67, 74, 84, 228, 301, 513 |
 SEQ ID NO : 19 | 9 | SEQ ID NO : 395 , SEQ ID NO : 396 , SEQ ID NO : 397 , SEQ ID NO : 398 | AFP, TRIM26, ZNF173 |
 15 3, 7, 149 |
 SEQ ID NO : 20 | 9 | SEQ ID NO : 399 , SEQ ID NO : 400 , SEQ ID NO : 401 , SEQ ID NO : 402 , SEQ ID NO : 403 , SEQ
 ID NO : 404 | RPA2 | 7, 65, 452, 545, 546 |
 SEQ ID NO : 21 | 9 | SEQ ID NO : 405 , SEQ ID NO : 406 , SEQ ID NO : 407 , SEQ ID NO : 408 , SEQ ID NO : 409 , SEQ
 ID NO : 410 , SEQ ID NO : 411 | CDC18L, CDC6, HSCDC6 | 8, 55, 56, 65, 67, 221, 529, 547, 548, 549, 550 |
 SEQ ID NO : 22 | 9 | SEQ ID NO : 412 , SEQ ID NO : 413 , SEQ ID NO : 414 , SEQ ID NO : 415 , SEQ ID NO : 416 , SEQ
 20 ID NO : 417 , SEQ ID NO : 418 , SEQ ID NO : 419 , SEQ ID NO : 420 , SEQ ID NO : 421 , SEQ ID NO : 422 | CBL, CBL2 ,
 CBLB | 23, 36, 89, 140, 149, 329 |
 SEQ ID NO : 23 | 9 | SEQ ID NO : 423 , SEQ ID NO : 424 , SEQ ID NO : 425 , SEQ ID NO : 426 , SEQ ID NO : 427 , SEQ
 ID NO : 428 , SEQ ID NO : 429 , SEQ ID NO : 430 , SEQ ID NO : 431 , SEQ ID NO : 432 | KIAA0898, MUL |
 25 SEQ ID NO : 24 | 7, 9 | SEQ ID NO : 433 , SEQ ID NO : 434 , SEQ ID NO : 435 , SEQ ID NO : 436 , SEQ ID NO : 437 ,
 SEQ ID NO : 438 , SEQ ID NO : 439 , SEQ ID NO : 440 , SEQ ID NO : 441 , SEQ ID NO : 442 , SEQ ID NO : 443 , SEQ
 ID NO : 444 , SEQ ID NO : 445 | PPP2R1A, PPP2R1B | 2, 137, 558, 559 |
 SEQ ID NO : 25 | 9 | SEQ ID NO : 446 , SEQ ID NO : 447 , SEQ ID NO : 448 , SEQ ID NO : 449 , SEQ ID NO : 450 , SEQ
 ID NO : 451 , SEQ ID NO : 452 , SEQ ID NO : 453 | HUMNDME, ME1 | 45, 110, 158, 303, 561, 562, 563 |
 SEQ ID NO : 26 | 9 | SEQ ID NO : 454 , SEQ ID NO : 455 , SEQ ID NO : 456 , SEQ ID NO : 457 , SEQ ID NO : 458 , SEQ
 30 ID NO : 459 , SEQ ID NO : 460 , SEQ ID NO : 461 , SEQ ID NO : 462 , SEQ ID NO : 463 | M11S1 | 31 |
 SEQ ID NO : 27 | 9 | SEQ ID NO : 464 , SEQ ID NO : 465 , SEQ ID NO : 466 , SEQ ID NO : 467 , SEQ ID NO : 468 , SEQ
 ID NO : 469 , SEQ ID NO : 470 , SEQ ID NO : 471 , SEQ ID NO : 472 , SEQ ID NO : 473 | BAL, BSSL, CEL, IRF4, LSIRF ,
 MUM1 | 7, 36, 54, 76, 568 |
 SEQ ID NO : 28 | 9 | SEQ ID NO : 474 , SEQ ID NO : 475 , SEQ ID NO : 476 , SEQ ID NO : 477 , SEQ ID NO : 478 , SEQ
 35 ID NO : 479 , SEQ ID NO : 480 , SEQ ID NO : 481 , SEQ ID NO : 482 , SEQ ID NO : 483 , SEQ ID NO : 484 , SEQ ID
 NO : 485 | MDH2 |
 SEQ ID NO : 29 | 9 | SEQ ID NO : 486 , SEQ ID NO : 487 , SEQ ID NO : 488 , SEQ ID NO : 489 , SEQ ID NO : 490 , SEQ

ID NO:491, SEQ ID NO:492, SEQ ID NO:493, SEQ ID NO:494, SEQ ID NO:495, SEQ ID NO:496, SEQ ID NO:497 | PCNA | 3, 7, 8, 56, 57, 60, 61, 62, 63, 64, 65, 66 |
 SEQ ID NO:30 | 9 | SEQ ID NO:498, SEQ ID NO:499, SEQ ID NO:500, SEQ ID NO:501, SEQ ID NO:502, SEQ ID NO:503, SEQ ID NO:504, SEQ ID NO:505, SEQ ID NO:506, SEQ ID NO:507, SEQ ID NO:508, SEQ ID NO:509, SEQ ID NO:510 | 4F2, 4F2HC, CD98, MD01, NACAE, SLC3A2 |
 5 SEQ ID NO:31 | 9 | SEQ ID NO:511, SEQ ID NO:512, SEQ ID NO:513, SEQ ID NO:514, SEQ ID NO:515, SEQ ID NO:516 | PKL.3 | 2 |
 SEQ ID NO:32 | 9 | SEQ ID NO:517, SEQ ID NO:518, SEQ ID NO:519, SEQ ID NO:520, SEQ ID NO:521, SEQ ID NO:522, SEQ ID NO:523, SEQ ID NO:524, SEQ ID NO:525, SEQ ID NO:526 | JNKK2, MAP2K7, MAPKK7,
 MKK7, PRKMK7 | 12, 89, 384, 579, 586 |
 10 SEQ ID NO:33 | 9 | SEQ ID NO:527, SEQ ID NO:528, SEQ ID NO:529, SEQ ID NO:530, SEQ ID NO:531, SEQ ID NO:532, SEQ ID NO:533, SEQ ID NO:534, SEQ ID NO:535, SEQ ID NO:536, SEQ ID NO:537, SEQ ID NO:538, SEQ ID NO:539, SEQ ID NO:540, SEQ ID NO:541, SEQ ID NO:542, SEQ ID NO:543, SEQ ID NO:544, SEQ ID NO:545, SEQ ID NO:546, SEQ ID NO:547 | MYC | 4, 8, 10, 17, 18, 23, 56, 57, 151, 371, 590 |
 SEQ ID NO:34 | 9 | SEQ ID NO:548, SEQ ID NO:549, SEQ ID NO:550, SEQ ID NO:551, SEQ ID NO:552, SEQ ID NO:553, SEQ ID NO:554 | DJ196E2 | 3, 2, HTATSF1, TAT-SF1 | 8, 10, 54, 151, 231, 556, 593 |
 SEQ ID NO:35 | 9, 13 | SEQ ID NO:555, SEQ ID NO:556, SEQ ID NO:557, SEQ ID NO:558, SEQ ID NO:559, SEQ ID NO:560, SEQ ID NO:561, SEQ ID NO:562, SEQ ID NO:563, SEQ ID NO:564, SEQ ID NO:565, SEQ ID NO:566, SEQ ID NO:567 | BRAF | 12, 87, 104, 176, 409 |
 15 SEQ ID NO:36 | 9 | SEQ ID NO:568, SEQ ID NO:569, SEQ ID NO:570, SEQ ID NO:571, SEQ ID NO:572, SEQ ID NO:573, SEQ ID NO:574 | DNAPK, DNPK1, HYRCK1, PRKDC, XRCC7 | 12, 60, 104, 129, 165, 166, 167 |
 SEQ ID NO:37 | 9, 13 | SEQ ID NO:575, SEQ ID NO:576, SEQ ID NO:577, SEQ ID NO:578, SEQ ID NO:579, SEQ ID NO:580, SEQ ID NO:581, SEQ ID NO:582 | EPS15R |
 SEQ ID NO:38 | 9, 24 | SEQ ID NO:583, SEQ ID NO:584, SEQ ID NO:585, SEQ ID NO:586, SEQ ID NO:587, SEQ ID NO:588 | HEC | 8, 35, 174, 601 |
 20 SEQ ID NO:39 | 9 | SEQ ID NO:589, SEQ ID NO:590, SEQ ID NO:591, SEQ ID NO:592, SEQ ID NO:593, SEQ ID NO:594 | 14.1, CD179B, IGI, IGL5, IGLJ1, IGLL1, IGO, IGVPB, POR1, VPREB1 | 47, 149, 161, 201, 306, 604, 605, 606 |
 SEQ ID NO:40 | 9 | SEQ ID NO:595, SEQ ID NO:596, SEQ ID NO:597, SEQ ID NO:598, SEQ ID NO:599, SEQ ID NO:600, SEQ ID NO:601, SEQ ID NO:602, SEQ ID NO:603, SEQ ID NO:604, SEQ ID NO:605, SEQ ID NO:606, SEQ ID NO:607, SEQ ID NO:608, SEQ ID NO:609, SEQ ID NO:610 | ENO1, ENO1L1, MBP-1, MPB1, NNE, PPH | 3, 7, 9, 10, 18, 23, 24, 25, 26 |
 25 SEQ ID NO:41 | 9 | SEQ ID NO:611, SEQ ID NO:612, SEQ ID NO:613, SEQ ID NO:614, SEQ ID NO:615, SEQ ID NO:616, SEQ ID NO:617, SEQ ID NO:618 | ITGB7 | 34, 186, 405, 408 |
 30 SEQ ID NO:42 | 9 | SEQ ID NO:619, SEQ ID NO:620, SEQ ID NO:621, SEQ ID NO:622, SEQ ID NO:623, SEQ ID NO:624, SEQ ID NO:625, SEQ ID NO:627, SEQ ID NO:628, SEQ ID NO:629, SEQ ID NO:630 | PCCB | 42, 71, 111, 615, 616 |

SEQ ID NO: 43 9 SEQ ID NO: 631, SEQ ID NO: 632, SEQ ID NO: 633, SEQ ID NO: 634, SEQ ID NO: 635, SEQ ID NO: 636, SEQ ID NO: 637, SEQ ID NO: 638, SEQ ID NO: 639, SEQ ID NO: 640, SEQ ID NO: 641, SEQ ID NO: 642, SEQ ID NO: 643, SEQ ID NO: 644, SEQ ID NO: 645, SEQ ID NO: 646, SEQ ID NO: 647, SEQ ID NO: 648 CBP, CREBBP, EP300, RSTS, RTS 8, 10, 14, 17, 23, 53, 67, 89, 106, 107, 118, 135, 332, 557 SEQ ID NO: 44 9 SEQ ID NO: 649, SEQ ID NO: 650, SEQ ID NO: 651, SEQ ID NO: 652, SEQ ID NO: 653 HGFR, MET 12, 31, 34, 36, 56, 57, 89, 631	5
SEQ ID NO: 45 9 SEQ ID NO: 654, SEQ ID NO: 655, SEQ ID NO: 656, SEQ ID NO: 657, SEQ ID NO: 658, SEQ ID NO: 659, SEQ ID NO: 660, SEQ ID NO: 661, SEQ ID NO: 662, SEQ ID NO: 663, SEQ ID NO: 664, SEQ ID NO: 665 CDHF12, HSCR1, MEN2A, MEN2B, MTC1, PTC, RET 12, 34, 36, 87, 89, 92, 107, 385, 448, 634	10
SEQ ID NO: 46 4 SEQ ID NO: 666, SEQ ID NO: 667, SEQ ID NO: 668, SEQ ID NO: 669, SEQ ID NO: 670, SEQ ID NO: 671, SEQ ID NO: 672, SEQ ID NO: 673 DEC-205, GP200-MR6, LY75 31, 34, 127, 129, 298, 385, 675 SEQ ID NO: 47 1 SEQ ID NO: 674, SEQ ID NO: 675, SEQ ID NO: 676, SEQ ID NO: 677, SEQ ID NO: 678, SEQ ID NO: 679, SEQ ID NO: 680, SEQ ID NO: 681 CD99, MIC2, MIC2X, MIC2Y 31, 67, 129, 154, 186, 187, 717 SEQ ID NO: 48 1 SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ ID NO: 687, SEQ ID NO: 688 CD71, TFR, TFRC 31, 331, 590, 720, 721, 722	15
SEQ ID NO: 49 1 SEQ ID NO: 689, SEQ ID NO: 690, SEQ ID NO: 691, SEQ ID NO: 692, SEQ ID NO: 693, SEQ ID NO: 694, SEQ ID NO: 695, SEQ ID NO: 696 CD49D, ITGA4 31, 34, 186, 408	20
SEQ ID NO: 50 10 SEQ ID NO: 256, SEQ ID NO: 257, SEQ ID NO: 258, SEQ ID NO: 259, SEQ ID NO: 260, SEQ ID NO: 261, SEQ ID NO: 262, SEQ ID NO: 263, SEQ ID NO: 264 CD27L, CD27LG, CD70, KI-24 ANTIGEN, TNFSF7 31, 56, 57, 89, 120, 179, 387, 388, 389	25
SEQ ID NO: 51 10 SEQ ID NO: 697, SEQ ID NO: 698, SEQ ID NO: 699, SEQ ID NO: 700, SEQ ID NO: 701, SEQ ID NO: 719, SEQ ID NO: 720 CD18, ITGB2, LAD, LCAMB, LFA-1, MF17 34, 129, 141, 186, 405, 408, 766	30
SEQ ID NO: 54 10 SEQ ID NO: 721, SEQ ID NO: 722, SEQ ID NO: 723, SEQ ID NO: 724, SEQ ID NO: 725, SEQ ID NO: 726, SEQ ID NO: 727, SEQ ID NO: 728, SEQ ID NO: 729, SEQ ID NO: 730, SEQ ID NO: 731, SEQ ID NO: 732, SEQ ID NO: 733, SEQ ID NO: 734, SEQ ID NO: 735, SEQ ID NO: 736, SEQ ID NO: 737, SEQ ID NO: 738, SEQ ID NO: 739, SEQ ID NO: 740, SEQ ID NO: 741, SEQ ID NO: 742, SEQ ID NO: 743, SEQ ID NO: 744, SEQ ID NO: 745, SEQ ID NO: 746, SEQ ID NO: 747, SEQ ID NO: 748, SEQ ID NO: 749, SEQ ID NO: 750, SEQ ID NO: 751, SEQ ID NO: 752, SEQ ID NO: 753, SEQ ID NO: 754 BB2, CD54, ICAM1 129, 154, 186, 187, 219, 231, 331, 385, 717, 778	35
SEQ ID NO: 55 10 SEQ ID NO: 755, SEQ ID NO: 756, SEQ ID NO: 757, SEQ ID NO: 758, SEQ ID NO: 759, SEQ ID NO: 760 DUT1, ROBO1 31, 106, 107, 186, 408, 798	
SEQ ID NO: 56 10 SEQ ID NO: 761, SEQ ID NO: 762, SEQ ID NO: 763, SEQ ID NO: 764, SEQ ID NO: 765, SEQ ID NO: 766	

5 ID NO: 766, SEQ ID NO: 767, SEQ ID NO: 768, SEQ ID NO: 769, SEQ ID NO: 770, SEQ ID NO: 771, SEQ ID NO: 772, SEQ ID NO: 773, SEQ ID NO: 774, SEQ ID NO: 775, SEQ ID NO: 776, SEQ ID NO: 777, SEQ ID NO: 778, SEQ ID NO: 779, SEQ ID NO: 780|VRL, VRL-1|22,29,31,741,742,743,744,745| SEQ ID NO: 57|11,12|SEQ ID NO: 781, SEQ ID NO: 782, SEQ ID NO: 783, SEQ ID NO: 784, SEQ ID NO: 785, TM7LNL|31,34,120,127,129,154,161,186,201,298,359,379,386,389,778|

10 SEQ ID NO: 58|11,12|SEQ ID NO: 792, SEQ ID NO: 793, SEQ ID NO: 794, SEQ ID NO: 795, SEQ ID NO: 796, SEQ ID NO: 797, SEQ ID NO: 798, SEQ ID NO: 799, SEQ ID NO: 800, SEQ ID NO: 801, SEQ ID NO: 802, SEQ ID NO: 803, SEQ ID NO: 804, SEQ ID NO: 805, SEQ ID NO: 806, SEQ ID NO: 807, SEQ ID NO: 808, SEQ ID NO: 809|FH, FHC, LDLR|31,76,89,154,219,331,464,510,778,825,826,827,828|

15 SEQ ID NO: 59|11,12|SEQ ID NO: 792, SEQ ID NO: 793, SEQ ID NO: 794, SEQ ID NO: 795, SEQ ID NO: 796, SEQ ID NO: 797, SEQ ID NO: 798, SEQ ID NO: 799, SEQ ID NO: 800, SEQ ID NO: 801, SEQ ID NO: 802, SEQ ID NO: 803, SEQ ID NO: 804, SEQ ID NO: 805, SEQ ID NO: 806, SEQ ID NO: 807, SEQ ID NO: 808, SEQ ID NO: 809|FH, FHC, LDLR|31,76,89,154,219,331,464,510,778,825,826,827,828|

20 SEQ ID NO: 60|11,12|SEQ ID NO: 810, SEQ ID NO: 811, SEQ ID NO: 812, SEQ ID NO: 813, SEQ ID NO: 814| GP250,LR11,LRP9,SORL1,SORLA|31,331,778,838,839|

25 SEQ ID NO: 61|11,12|SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ ID NO: 687, SEQ ID NO: 688|CD71,TFR,TFRC|31,331,590,720,721,722|

30 SEQ ID NO: 62|11,12|SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ ID NO: 687, SEQ ID NO: 688|CD71,TFR,TFRC|31,331,590,720,721,722|

35 SEQ ID NO: 63|11,12|SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ ID NO: 687, SEQ ID NO: 688|CD71,TFR,TFRC|31,331,590,720,721,722|

NO:873, SEQ ID NO:874 | KIAA0343, NRCAM|2,31,107,186,187 |
 SEQ ID NO:69|5 |SEQ ID NO:875, SEQ ID NO:876, SEQ ID NO:877, SEQ ID NO:878, SEQ ID NO:879, SEQ
 ID NO:880, SEQ ID NO:881, SEQ ID NO:882, SEQ ID NO:883, SEQ ID NO:884 | CDD122, IL2RB|31,34,89,
 127,129,154,332,402 |

5 SEQ ID NO:70|5 |SEQ ID NO:885, SEQ ID NO:886, SEQ ID NO:887, SEQ ID NO:888, SEQ ID NO:889, SEQ
 ID NO:890, SEQ ID NO:891, SEQ ID NO:892, SEQ ID NO:893, SEQ ID NO:894, SEQ ID NO:895, SEQ ID
 NO:896, SEQ ID NO:897, SEQ ID NO:898, SEQ ID NO:899, SEQ ID NO:900, SEQ ID NO:901, SEQ ID
 NO:902, SEQ ID NO:903, SEQ ID NO:904 | CD23, CD23A, FCER2, SST|28,31,33,34,55,56,102,120,
 129,179,253,359,386,389,438,765,848,849,850,851 |

10 SEQ ID NO:71|5 |SEQ ID NO:885, SEQ ID NO:886, SEQ ID NO:887, SEQ ID NO:888, SEQ ID NO:889, SEQ
 ID NO:890, SEQ ID NO:891, SEQ ID NO:892, SEQ ID NO:893, SEQ ID NO:894, SEQ ID NO:895, SEQ ID
 NO:896, SEQ ID NO:897, SEQ ID NO:898, SEQ ID NO:899, SEQ ID NO:900, SEQ ID NO:901, SEQ ID
 NO:902, SEQ ID NO:903, SEQ ID NO:904 | CD23, CD23A, FCER2, SST|28,31,33,34,55,56,102,120,
 129,179,253,359,386,389,438,765,848,849,850,851 |

15 SEQ ID NO:72|5 |SEQ ID NO:885, SEQ ID NO:886, SEQ ID NO:887, SEQ ID NO:888, SEQ ID NO:889, SEQ
 ID NO:890, SEQ ID NO:891, SEQ ID NO:892, SEQ ID NO:893, SEQ ID NO:894, SEQ ID NO:895, SEQ ID
 NO:896, SEQ ID NO:897, SEQ ID NO:898, SEQ ID NO:899, SEQ ID NO:900, SEQ ID NO:901, SEQ ID
 NO:902, SEQ ID NO:903, SEQ ID NO:904 | CD23, CD23A, FCER2, SST|28,31,33,34,55,56,102,120,
 129,179,253,359,386,389,438,765,848,849,850,851 |

20 SEQ ID NO:73|5 |SEQ ID NO:885, SEQ ID NO:886, SEQ ID NO:887, SEQ ID NO:888, SEQ ID NO:889, SEQ
 ID NO:890, SEQ ID NO:891, SEQ ID NO:892, SEQ ID NO:893, SEQ ID NO:894, SEQ ID NO:895, SEQ ID
 NO:896, SEQ ID NO:897, SEQ ID NO:898, SEQ ID NO:899, SEQ ID NO:900, SEQ ID NO:901, SEQ ID
 NO:902, SEQ ID NO:903, SEQ ID NO:904 | CD23, CD23A, FCER2, SST|28,31,33,34,55,56,102,120,
 129,179,253,359,386,389,438,765,848,849,850,851 |

25 SEQ ID NO:74|5 |SEQ ID NO:885, SEQ ID NO:886, SEQ ID NO:887, SEQ ID NO:888, SEQ ID NO:889, SEQ
 ID NO:890, SEQ ID NO:891, SEQ ID NO:892, SEQ ID NO:893, SEQ ID NO:894, SEQ ID NO:895, SEQ ID
 NO:896, SEQ ID NO:897, SEQ ID NO:898, SEQ ID NO:899, SEQ ID NO:900, SEQ ID NO:901, SEQ ID
 NO:902, SEQ ID NO:903, SEQ ID NO:904 | CD23, CD23A, FCER2, SST|28,31,33,34,55,56,102,120,
 129,179,253,359,386,389,438,765,848,849,850,851 |

30 SEQ ID NO:75|5 |SEQ ID NO:905, SEQ ID NO:906, SEQ ID NO:907, SEQ ID NO:908, SEQ ID NO:909, SEQ
 ID NO:910 | ADAM9, KIAA0021, MCMP, MDC9, MLTNG|31,47,51,212,640,765 |
 SEQ ID NO:76|5 |SEQ ID NO:911, SEQ ID NO:912, SEQ ID NO:913, SEQ ID NO:914, SEQ ID NO:915, SEQ
 ID NO:916, SEQ ID NO:917, SEQ ID NO:918, SEQ ID NO:919, SEQ ID NO:920 | HPTPE, PTPRE|31,34,
 47,67,68,468,858 |

35 SEQ ID NO:77|5 |SEQ ID NO:682, SEQ ID NO:683, SEQ ID NO:684, SEQ ID NO:685, SEQ ID NO:686, SEQ
 ID NO:687, SEQ ID NO:688 | CD71, TFR, TERC|31,331,590,720,721,722 |
 SEQ ID NO:78|5 |SEQ ID NO:682, SEQ ID NO:683, SEQ ID NO:684, SEQ ID NO:685, SEQ ID NO:921, SEQ

ID NO: 686, SEQ ID NO: 687, SEQ ID NO: 688 | CD71, TFR, TFRC | 31, 331, 590, 720, 721, 722 |
 SEQ ID NO: 79 | 5 | SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ
 ID NO: 687, SEQ ID NO: 688 | CD71, TFR, TFRC | 31, 331, 590, 720, 721, 722 |
 SEQ ID NO: 80 | 5 | SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ
 ID NO: 687, SEQ ID NO: 688 | CD71, TFR, TFRC | 31, 331, 590, 720, 721, 722 |
 SEQ ID NO: 81 | 5 | SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ
 ID NO: 687, SEQ ID NO: 688 | CD71, TFR, TFRC | 31, 331, 590, 720, 721, 722 |
 SEQ ID NO: 82 | 5 | SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ
 ID NO: 686, SEQ ID NO: 687, SEQ ID NO: 688 | CD71, TFR, TFRC | 31, 331, 590, 720, 721, 722 |
 SEQ ID NO: 83 | 5 | SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ
 ID NO: 687, SEQ ID NO: 688 | CD71, TFR, TFRC | 31, 331, 590, 720, 721, 722 |
 SEQ ID NO: 84 | 5 | SEQ ID NO: 682, SEQ ID NO: 683, SEQ ID NO: 684, SEQ ID NO: 685, SEQ ID NO: 686, SEQ
 ID NO: 687, SEQ ID NO: 688 | CD71, TFR, TFRC | 31, 331, 590, 720, 721, 722 |
 SEQ ID NO: 85 | 1, 13 | SEQ ID NO: 923, SEQ ID NO: 924, SEQ ID NO: 925, SEQ ID NO: 926, SEQ ID NO: 927,
 SEQ ID NO: 928 | PRKAG1 | 12, 87, 89, 565, 869 |
 SEQ ID NO: 86 | 4 | CRL1, TCCR, WSX-1, WSX1 | 31, 34, 127, 129, 386, 778 |
 SEQ ID NO: 87 | 1 | SEQ ID NO: 256, SEQ ID NO: 258, SEQ ID NO: 262, SEQ ID NO: 264 | CD27LG, CD70,
 KI-24 ANTIGEN, TMFSF7 | 31, 56, 57, 89, 120, 179, 387, 388, 389 |
 SEQ ID NO: 88 | 18 | SEQ ID NO: 929, SEQ ID NO: 930, SEQ ID NO: 931, SEQ ID NO: 932, SEQ ID NO: 933, SEQ
 ID NO: 934, SEQ ID NO: 935, SEQ ID NO: 936, SEQ ID NO: 937, SEQ ID NO: 938, SEQ ID NO: 939 | CD49F,
 ITGA6 | 34, 200, 224, 405, 408, 923 |
 SEQ ID NO: 89 | 18 | SEQ ID NO: 929, SEQ ID NO: 930, SEQ ID NO: 931, SEQ ID NO: 933, SEQ ID NO: 935, SEQ
 ID NO: 936, SEQ ID NO: 937, SEQ ID NO: 938, SEQ ID NO: 939 | CD49F, ITGA6 | 34, 200, 224, 405, 408, 923 |
 SEQ ID NO: 90 | 18 | SEQ ID NO: 940, SEQ ID NO: 941, SEQ ID NO: 942, SEQ ID NO: 943, SEQ ID NO: 944, SEQ
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 NO: 975, SEQ ID NO: 976, SEQ ID NO: 977 | CALM, CLTH, DJ167A19.1, FILT4, KIAA0556, PCL, PICALM, SNAP91,
 VEGFR3 | 12, 31, 34, 135, 332, 411, 461, 639, 815 |
 SEQ ID NO: 91 | 19 | SEQ ID NO: 978, SEQ ID NO: 979, SEQ ID NO: 980, SEQ ID NO: 981, SEQ ID NO: 982, SEQ
 ID NO: 983, SEQ ID NO: 984, SEQ ID NO: 985, SEQ ID NO: 986, SEQ ID NO: 987 | CD222, CIMPR, IGF2R, M6P-R,
 MPRI | 2, 31, 34, 89, 219, 331, 385, 839, 932 |
 SEQ ID NO: 92 | 19 | SEQ ID NO: 846, SEQ ID NO: 847, SEQ ID NO: 848, SEQ ID NO: 849, SEQ ID NO: 850, SEQ
 ID NO: 851, SEQ ID NO: 852, SEQ ID NO: 853, SEQ ID NO: 854, SEQ ID NO: 855, SEQ ID NO: 856, SEQ ID
 NO: 857, SEQ ID NO: 858, SEQ ID NO: 859, SEQ ID NO: 860, SEQ ID NO: 861 | IFNGR, IFNGR1 | 31, 34, 89, 127,

129, 154, 229, 230, 843, 844 |
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SEQ ID NO:94|20|SEQ ID NO:792, SEQ ID NO:793, SEQ ID NO:794, SEQ ID NO:1006, SEQ ID NO:795, SEQ ID NO:796, SEQ ID NO:797, SEQ ID NO:798, SEQ ID NO:799, SEQ ID NO:800, SEQ ID NO:801, SEQ ID NO:802, SEQ ID NO:803, SEQ ID NO:804, SEQ ID NO:805, SEQ ID NO:806, SEQ ID NO:807, SEQ ID NO:808, SEQ ID NO:809|FH, FHC, LDLR|31, 76, 89, 154, 219, 331, 464, 510, 778, 825, 826,
10 827, 828 |
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15 31, 34, 148, 188, 553 |
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44, 110, 147, 647, 700, 952 |
25 SEQ ID NO:98|7, 24|SEQ ID NO:1055, SEQ ID NO:1056, SEQ ID NO:1057, SEQ ID NO:1058, SEQ ID NO:1059, SEQ ID NO:1060, SEQ ID NO:1061, SEQ ID NO:1062, SEQ ID NO:1063|PLXN#, PLXNC1, PLXNC1,
VESPR|129, 154, 186, 209, 219, 231, 331, 387 |
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30 8, 56, 57, 60, 61, 62, 63, 64, 65, 66 |
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956 |
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 NO:1336, SEQ ID NO:1337, SEQ ID NO:1338, SEQ ID NO:1339|37LRP, LAMBR, LAMR1, LRP, P40, RPSA|31, 34,
 36, 72, 162, 186, 352, 386, 395, 396, 397|
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 154, 635, 1012|
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 777, 1035, 1036|
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 CRACCI, SMARCC1, SRG3|10, 118, 151, 189, 199|
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 NO:1381, SEQ ID NO:1382, SEQ ID NO:1383, SEQ ID NO:1384, SEQ ID NO:1385, SEQ ID NO:1386|PDK1,
 PDPK1|12, 87, 104, 105, 133, 512, 1044|
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 SEQ ID NO:120|13, 24|SEQ ID NO:1401, SEQ ID NO:1402, SEQ ID NO:1403, SEQ ID NO:1404, SEQ ID NO:1405, SEQ ID NO:1406, SEQ ID NO:1407, SEQ ID NO:1408, SEQ ID NO:1409, SEQ ID NO:1410, SEQ ID NO:1411, SEQ ID NO:1412, SEQ ID NO:1413, SEQ ID NO:1414, SEQ ID NO:1415, SEQ ID NO:1416|ISGF-3,
 STAT1, STAT91|3, 7, 8, 10, 16, 23, 53, 62, 67, 89, 179, 263, 265, 781, 782, 783, 784, 785, 786|
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 GPR34|31, 34, 359, 379|
 SEQ ID NO:122|13|SEQ ID NO:1439, SEQ ID NO:1440, SEQ ID NO:1441, SEQ ID NO:1442, SEQ ID NO:1443|HTOLL, TLR4|31, 34, 89, 127, 129, 385, 961|
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 P40, RPSA|31, 34, 36, 72, 162, 186, 352, 386, 395, 396, 397|
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 154, 635, 1012|
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 CEACAM6, CEACAM8, CGM6, NCA, NCA-95|31, 34, 89, 120, 127, 129, 154, 297, 389, 438|
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GP250, LR11, LRP9, SORL1, SORL1A|31, 331, 778, 838, 839|

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MS4A1A, MS4A2|22, 31, 57, 129, 359, 789, 1110|

15 SEQ ID NO:131|25, 26|SEQ ID NO:1515, SEQ ID NO:1516, SEQ ID NO:1517, SEQ ID NO:1518, SEQ ID NO:1519, SEQ ID NO:1520, SEQ ID NO:1521, SEQ ID NO:1522, SEQ ID NO:1523, SEQ ID NO:1524, SEQ ID NO:1525, SEQ ID NO:1526, SEQ ID NO:1527, SEQ ID NO:1528, SEQ ID NO:1529, SEQ ID NO:1530, SEQ ID NO:1531, SEQ ID NO:1532, SEQ ID NO:1533, SEQ ID NO:1534, SEQ ID NO:1535|B1, BP35, CD20, MS4A1,
MS4A1A, MS4A2|22, 31, 57, 129, 359, 789, 1110|

20 SEQ ID NO:132|25, 26|SEQ ID NO:1515, SEQ ID NO:1516, SEQ ID NO:1517, SEQ ID NO:1518, SEQ ID NO:1519, SEQ ID NO:1520, SEQ ID NO:1521, SEQ ID NO:1522, SEQ ID NO:1523, SEQ ID NO:1524, SEQ ID NO:1525, SEQ ID NO:1526, SEQ ID NO:1527, SEQ ID NO:1528, SEQ ID NO:1529, SEQ ID NO:1530, SEQ ID NO:1531, SEQ ID NO:1532, SEQ ID NO:1533, SEQ ID NO:1534, SEQ ID NO:1535|B1, BP35, CD20, MS4A1,
MS4A1A, MS4A2|22, 31, 57, 129, 359, 789, 1110|

25 SEQ ID NO:133|25, 26|SEQ ID NO:682, SEQ ID NO:683, SEQ ID NO:684, SEQ ID NO:685, SEQ ID NO:686,
SEQ ID NO:687, SEQ ID NO:688|CD71, TFR, TFRC|31, 331, 590, 720, 721, 722|
SEQ ID NO:134|28|SEQ ID NO:1536, SEQ ID NO:1537, SEQ ID NO:1538, SEQ ID NO:1539, SEQ ID NO:1540|WNT5A|36, 68, 89, 91, 120, 345, 387, 389, 438, 1114|
SEQ ID NO:135|28|SEQ ID NO:1541, SEQ ID NO:1542, SEQ ID NO:1543, SEQ ID NO:1544|CB2, CNR2|31,
34, 127, 129, 154, 754, 1116, 1117, 1118|

30 SEQ ID NO:136|28|SEQ ID NO:1545, SEQ ID NO:1546, SEQ ID NO:1547, SEQ ID NO:1548, SEQ ID NO:1549, SEQ ID NO:1550, SEQ ID NO:1551, SEQ ID NO:1552, SEQ ID NO:1553, SEQ ID NO:1554|RRM1|
SEQ ID NO:137|24|SEQ ID NO:486, SEQ ID NO:487, SEQ ID NO:488, SEQ ID NO:489, SEQ ID NO:490,
SEQ ID NO:1064, SEQ ID NO:491, SEQ ID NO:494, SEQ ID NO:495, SEQ ID NO:496, SEQ ID NO:497|PCNA|
3, 7, 8, 56, 57, 60, 61, 62, 63, 64, 65, 66|

35 SEQ ID NO:138|24|SEQ ID NO:1401, SEQ ID NO:1402, SEQ ID NO:1403, SEQ ID NO:1555, SEQ ID NO:1405, SEQ ID NO:1406, SEQ ID NO:1407, SEQ ID NO:1408, SEQ ID NO:1410, SEQ ID NO:1411, SEQ ID NO:1412, SEQ ID NO:1413, SEQ ID NO:1414, SEQ ID NO:1415, SEQ ID NO:1416|ISGF-3,

STAT1, STAT91|3, 7, 8, 10, 16, 23, 53, 62, 67, 89, 179, 263, 265, 781, 782, 783, 784, 785, 786|
 SEQ ID NO:139|24|SEQ ID NO:1556, SEQ ID NO:1557, SEQ ID NO:1558, SEQ ID NO:1559, SEQ ID
 NO:1560|PCM1|67, 134, 501, 1121|
 SEQ ID NO:140|24|SEQ ID NO:1561, SEQ ID NO:1562, SEQ ID NO:1563, SEQ ID NO:1564, SEQ ID
 NO:1565, SEQ ID NO:1566, SEQ ID NO:1567, SEQ ID NO:1568, SEQ ID NO:1569|CADTK, CAK BETA, CAKB,
 FAK2, PKB, PTK, PTK2B, PYRK2, RAFTK|12, 56, 67, 87, 88, 89, 179, 186, 332, 344, 388, 448, 579, 1122, 1123|
 SEQ ID NO:141|1|SEQ ID NO:1570, SEQ ID NO:1571, SEQ ID NO:1572, SEQ ID NO:1573, SEQ ID
 NO:1574, SEQ ID NO:1575, SEQ ID NO:1576, SEQ ID NO:1577|TETRAN, TETTRAN|29, 209, 711, 712, 713|
 SEQ ID NO:142|13|SEQ ID NO:1578, SEQ ID NO:1579, SEQ ID NO:1580, SEQ ID NO:1581, SEQ ID
 NO:1582, SEQ ID NO:1583, SEQ ID NO:1584, SEQ ID NO:1585, SEQ ID NO:1586, SEQ ID NO:1587, SEQ ID
 NO:1588, SEQ ID NO:1589, SEQ ID NO:1590, SEQ ID NO:1591, SEQ ID NO:1592, SEQ ID NO:1593, SEQ ID
 NO:1594, SEQ ID NO:1595, SEQ ID NO:1596, SEQ ID NO:1597, SEQ ID NO:1598, SEQ ID NO:1599, SEQ ID
 NO:1600, SEQ ID NO:1601, SEQ ID NO:1602, SEQ ID NO:1603|MT-SP2, TMPRSS3, TMPRSS4|
 SEQ ID NO:143|15|SEQ ID NO:929, SEQ ID NO:930, SEQ ID NO:931, SEQ ID NO:932, SEQ ID NO:933,
 SEQ ID NO:1604, SEQ ID NO:934, SEQ ID NO:935, SEQ ID NO:936, SEQ ID NO:937, SEQ ID NO:938, SEQ
 ID NO:939|CD49F ITGA6|34, 200, 224, 405, 408, 923|
 SEQ ID NO:144|18|SEQ ID NO:1605, SEQ ID NO:1606, SEQ ID NO:1607, SEQ ID NO:1608, SEQ ID
 NO:1609, SEQ ID NO:1610, SEQ ID NO:1611, SEQ ID NO:1612, SEQ ID NO:1613, SEQ ID NO:1614, SEQ ID
 NO:1615, SEQ ID NO:1616, SEQ ID NO:1617, SEQ ID NO:1618, SEQ ID NO:1619, SEQ ID NO:1620, SEQ ID
 NO:1621, SEQ ID NO:1622, SEQ ID NO:1623, SEQ ID NO:1624, SEQ ID NO:1625, SEQ ID NO:1626, SEQ ID
 NO:1627, SEQ ID NO:1628, SEQ ID NO:1629, SEQ ID NO:1630, SEQ ID NO:1631, SEQ ID NO:1632, SEQ ID
 NO:1633, SEQ ID NO:1634, SEQ ID NO:1635, SEQ ID NO:1636, SEQ ID NO:1637, SEQ ID NO:1638, SEQ ID
 NO:1639, SEQ ID NO:1640|B-ALPHA-1, K-ALPHA-1, MGC10851, MGC14580, TUBA3, TUBA6|106, 132, 134, 163,
 319, 320, 321|
 SEQ ID NO:145|28|SEQ ID NO:1641, SEQ ID NO:1642, SEQ ID NO:1643, SEQ ID NO:1644, SEQ ID
 NO:1645, SEQ ID NO:1646, SEQ ID NO:1647|CD49C, GAPB3, ITGA3, VL3A|34, 200, 405, 408|
 SEQ ID NO:146|28|SEQ ID NO:1648, SEQ ID NO:1649, SEQ ID NO:1650, SEQ ID NO:1651, SEQ ID
 NO:1652, SEQ ID NO:1653, SEQ ID NO:1654, SEQ ID NO:1655, SEQ ID NO:1656, SEQ ID NO:1657, SEQ ID
 NO:1658, SEQ ID NO:1659|CMD1A, EMD2, FPLD, LDB1, LFP, LMN1, LMNA|132, 134, 320, 608, 627, 628|
 SEQ ID NO:147|24|SEQ ID NO:689, SEQ ID NO:1660, SEQ ID NO:690, SEQ ID NO:691, SEQ ID NO:692,
 SEQ ID NO:693, SEQ ID NO:694, SEQ ID NO:695, SEQ ID NO:696, SEQ ID NO:1661|CD49D, ITGA4|31, 34,
 186, 408|
 SEQ ID NO:148|24|SEQ ID NO:1662, SEQ ID NO:1663, SEQ ID NO:1664, SEQ ID NO:1665, SEQ ID
 NO:1666, SEQ ID NO:1667, SEQ ID NO:1668, SEQ ID NO:1669, SEQ ID NO:1670, SEQ ID NO:1671, SEQ ID
 NO:1672, SEQ ID NO:1673|B29, CD79B, IGB|31, 89, 127, 129|
 35 SEQ ID NO:149|24|SEQ ID NO:486, SEQ ID NO:487, SEQ ID NO:488, SEQ ID NO:489, SEQ ID NO:490,
 SEQ ID NO:491, SEQ ID NO:493, SEQ ID NO:494, SEQ ID NO:495, SEQ ID NO:496, SEQ ID NO:497|PCNA|

NO:1489,SEQ ID NO:1490,SEQ ID NO:1491,SEQ ID NO:1492,SEQ ID NO:1493,SEQ ID NO:1495,SEQ ID
 NO:1496,SEQ ID NO:1497,SEQ ID NO:1500,SEQ ID NO:1501,SEQ ID NO:1502,SEQ ID NO:1503,SEQ ID
 NO:1504,SEQ ID NO:1505,SEQ ID NO:1506,SEQ ID NO:1507,SEQ ID NO:1245|CD66B,CD66C,CD66E,
 CD67,CEA,CEACAM5,CEACAM6,CEACAM8,CGM6,NCA,NCA-95|31,34,89,120,127,129,154,297,389,438|
 SEQ ID NO:161|8,15|SEQ ID NO:1320,SEQ ID NO:1321,SEQ ID NO:1322,SEQ ID NO:1323,SEQ ID
 NO:1325,SEQ ID NO:1326,SEQ ID NO:1327,SEQ ID NO:1328,SEQ ID NO:1329,SEQ ID NO:1330,SEQ ID
 NO:1331,SEQ ID NO:1332,SEQ ID NO:1333,SEQ ID NO:1336,SEQ ID NO:1337,SEQ ID NO:1338|37LRP,
 LAMBR,LAMR1,LRP,P40,RPSA|31,34,36,72,162,186,352,386,395,396,397|
 SEQ ID NO:162|15|SEQ ID NO:1702,SEQ ID NO:1703,SEQ ID NO:1704,SEQ ID NO:1705,SEQ ID
 NO:1706,SEQ ID NO:1707,SEQ ID NO:1708,SEQ ID NO:1709,SEQ ID NO:1710,SEQ ID NO:1711,SEQ ID
 NO:1712,SEQ ID NO:1713|5F7,BSG,CD147,OK|89,242,386,739|
 SEQ ID NO:163|15|SEQ ID NO:988,SEQ ID NO:989,SEQ ID NO:990,SEQ ID NO:991,SEQ ID NO:992,
 SEQ ID NO:993,SEQ ID NO:994,SEQ ID NO:995,SEQ ID NO:996,SEQ ID NO:998,SEQ ID NO:999,SEQ
 ID NO:1000,SEQ ID NO:1001,SEQ ID NO:1002,SEQ ID NO:1003,SEQ ID NO:1004,SEQ ID NO:1005|
 CALLA,CD10,MME|31,36,47,49,51,120,154,282,389,640|
 SEQ ID NO:164|15|SEQ ID NO:1321,SEQ ID NO:1322,SEQ ID NO:1323,SEQ ID NO:1324,SEQ ID
 NO:1325,SEQ ID NO:1326,SEQ ID NO:1327,SEQ ID NO:1328,SEQ ID NO:1329,SEQ ID NO:1330,SEQ ID
 NO:1331,SEQ ID NO:1332,SEQ ID NO:1333,SEQ ID NO:1444,SEQ ID NO:1445,SEQ ID NO:1446,SEQ ID
 NO:1447,SEQ ID NO:1336,SEQ ID NO:1337,SEQ ID NO:1338,SEQ ID NO:1339|37LRP,LAMBR,LAMR1,LRP,
 P40,RPSA|31,34,36,72,162,186,352,386,395,396,397|
 SEQ ID NO:165|15|SEQ ID NO:1714,SEQ ID NO:1715,SEQ ID NO:1716,SEQ ID NO:1717,SEQ ID
 NO:1718,SEQ ID NO:1719,SEQ ID NO:1720,SEQ ID NO:1721,SEQ ID NO:1722,SEQ ID NO:1723,SEQ ID
 NO:1724,SEQ ID NO:1725,SEQ ID NO:1726|BA2,CD9,MIC3,MRX50,P24|31,129,154|
 SEQ ID NO:166|15|SEQ ID NO:1321,SEQ ID NO:1322,SEQ ID NO:1323,SEQ ID NO:1324,SEQ ID
 NO:1325,SEQ ID NO:1326,SEQ ID NO:1327,SEQ ID NO:1328,SEQ ID NO:1329,SEQ ID NO:1330,SEQ ID
 NO:1331,SEQ ID NO:1332,SEQ ID NO:1333,SEQ ID NO:1444,SEQ ID NO:1445,SEQ ID NO:1446,SEQ ID
 NO:1447,SEQ ID NO:1336,SEQ ID NO:1337,SEQ ID NO:1338,SEQ ID NO:1339|37LRP,LAMBR,LAMR1,LRP,
 P40,RPSA|31,34,36,72,162,186,352,386,395,396,397|
 SEQ ID NO:167|15|SEQ ID NO:1727,SEQ ID NO:1728,SEQ ID NO:1729,SEQ ID NO:1730,SEQ ID
 NO:1731,SEQ ID NO:1732,SEQ ID NO:1733,SEQ ID NO:1734|HPR6,6,MPR,PGRMC1|31,34,1180|
 SEQ ID NO:168|13|SEQ ID NO:1735,SEQ ID NO:1736,SEQ ID NO:1737,SEQ ID NO:1738,SEQ ID
 NO:1739,SEQ ID NO:1740,SEQ ID NO:1741,SEQ ID NO:1742,SEQ ID NO:1743,SEQ ID NO:1744|CD11A,
 ITGAL,LFA1-LFA1A|34,161,186,201,405,408|
 SEQ ID NO:169|13|SEQ ID NO:1745,SEQ ID NO:1746,SEQ ID NO:1747,SEQ ID NO:1748,SEQ ID
 NO:1749,SEQ ID NO:1750,SEQ ID NO:1751,SEQ ID NO:1752,SEQ ID NO:1753|ADAM17,CD156B,CSV_P,
 TACE|31,47,51,120,242,389,640|
 SEQ ID NO:170|15|SEQ ID NO:682,SEQ ID NO:683,SEQ ID NO:684,SEQ ID NO:685,SEQ ID NO:686,

SEQ ID NO: 687, SEQ ID NO: 688 | CD71, TFR, TFRC | 31, 331, 590, 720, 721, 722 |
 SEQ ID NO: 171 | 15 | SEQ ID NO: 978, SEQ ID NO: 980, SEQ ID NO: 981, SEQ ID NO: 982,
 SEQ ID NO: 983, SEQ ID NO: 985, SEQ ID NO: 986, SEQ ID NO: 987 | CD222, C1MPR, IGF2R, M6P-R, MPRI | 2, 31,
 34, 89, 219, 331, 385, 839, 932 |

5 SEQ ID NO: 172 | 15 | SEQ ID NO: 1320, SEQ ID NO: 1321, SEQ ID NO: 1322, SEQ ID NO: 1323, SEQ ID
 NO: 1324, SEQ ID NO: 1325, SEQ ID NO: 1326, SEQ ID NO: 1327, SEQ ID NO: 1328, SEQ ID NO: 1329, SEQ ID
 NO: 1330, SEQ ID NO: 1331, SEQ ID NO: 1332, SEQ ID NO: 1333, SEQ ID NO: 1334, SEQ ID NO: 1335, SEQ ID
 NO: 1754, SEQ ID NO: 1336, SEQ ID NO: 1337, SEQ ID NO: 1338, SEQ ID NO: 1339 | 37LRP, LAMBR, LAMR1, LRP,
 P40, RPSA | 31, 34, 36, 72, 162, 186, 352, 386, 395, 396, 397 |

10 SEQ ID NO: 173 | 8, 15 | SEQ ID NO: 1321, SEQ ID NO: 1322, SEQ ID NO: 1323, SEQ ID NO: 1324, SEQ ID
 NO: 1325, SEQ ID NO: 1326, SEQ ID NO: 1327, SEQ ID NO: 1328, SEQ ID NO: 1329, SEQ ID NO: 1330, SEQ ID
 NO: 1331, SEQ ID NO: 1332, SEQ ID NO: 1333, SEQ ID NO: 1444, SEQ ID NO: 1445, SEQ ID NO: 1446, SEQ ID
 NO: 1447, SEQ ID NO: 1336, SEQ ID NO: 1337, SEQ ID NO: 1338, SEQ ID NO: 1339 | 37LRP, LAMBR, LAMR1, LRP,
 P40, RPSA | 31, 34, 36, 72, 162, 186, 352, 386, 395, 396, 397 |

15 SEQ ID NO: 174 | 8, 15 | SEQ ID NO: 1321, SEQ ID NO: 1322, SEQ ID NO: 1323, SEQ ID NO: 1324, SEQ ID
 NO: 1325, SEQ ID NO: 1326, SEQ ID NO: 1327, SEQ ID NO: 1328, SEQ ID NO: 1330, SEQ ID NO: 1331, SEQ ID
 NO: 1332, SEQ ID NO: 1755, SEQ ID NO: 1336, SEQ ID NO: 1337, SEQ ID NO: 1338, SEQ ID NO: 1339 | 37LRP,
 LAMBR, LAMR1, LRP, P40, RPSA | 31, 34, 36, 72, 162, 186, 352, 386, 395, 396, 397 |

20 SEQ ID NO: 175 | 13 | SEQ ID NO: 1756, SEQ ID NO: 1757, SEQ ID NO: 1758, SEQ ID NO: 1759, SEQ ID
 NO: 1760, SEQ ID NO: 1761, SEQ ID NO: 1762, SEQ ID NO: 1763, SEQ ID NO: 1764, SEQ ID NO: 1765, SEQ ID
 NO: 1766, SEQ ID NO: 1767, SEQ ID NO: 1768, SEQ ID NO: 1769, SEQ ID NO: 1770, SEQ ID NO: 1771, SEQ ID
 NO: 1772, SEQ ID NO: 1773 | IFNAR, IFNAR1, IFRC | 31, 34, 129, 229, 263, 386, 1247 |

25 SEQ ID NO: 176 | 13 | SEQ ID NO: 929, SEQ ID NO: 930, SEQ ID NO: 931, SEQ ID NO: 932, SEQ ID NO: 933,
 SEQ ID NO: 1774, SEQ ID NO: 934, SEQ ID NO: 935, SEQ ID NO: 936, SEQ ID NO: 937, SEQ ID NO: 938, SEQ
 ID NO: 939 | CD49F, ITGA6 | 34, 200, 224, 405, 408, 923 |

30 SEQ ID NO: 177 | 13 | SEQ ID NO: 1780, SEQ ID NO: 1781, SEQ ID NO: 1777, SEQ ID NO: 1778, SEQ ID
 NO: 1779, SEQ ID NO: 1782, SEQ ID NO: 1783, SEQ ID NO: 1784 | CTNNB1,
 CTNNB1 | 1, 2, 8, 10, 16, 36, 89, 151, 154, 186, 790 |

35 SEQ ID NO: 178 | 13 | SEQ ID NO: 1785, SEQ ID NO: 1786, SEQ ID NO: 1787, SEQ ID NO: 1788, SEQ ID
 NO: 1789, SEQ ID NO: 1790, SEQ ID NO: 1791, SEQ ID NO: 1792, SEQ ID NO: 1793, SEQ ID NO: 1794, SEQ ID
 NO: 1795, SEQ ID NO: 1796 | CD98, D16S469E, E16, LAT1, MPE16, SLC7A5 | 29, 111, 154, 712, 77, 1189 |
 SEQ ID NO: 179 | 13 | SEQ ID NO: 1797, SEQ ID NO: 1798, SEQ ID NO: 1799, SEQ ID NO: 1800, SEQ ID
 NO: 1801, SEQ ID NO: 1802 | NKTR | 129, 154, 254, 255, 773 |

40 SEQ ID NO: 180 | 13 | SEQ ID NO: 1803, SEQ ID NO: 1804, SEQ ID NO: 1805, SEQ ID NO: 1806, SEQ ID
 NO: 1807, SEQ ID NO: 1808, SEQ ID NO: 1810, SEQ ID NO: 1811, SEQ ID NO: 1812, SEQ ID
 NO: 1813, SEQ ID NO: 1814 | DCR3, DJ583P15.1.1, DKFZP434C013, KIAA1088, M68, NHL, TNFRSF6B, TR6 | 18, 34,
 36, 68, 81, 138, 176, 179, 385 |

SEQ ID NO:181|13|SEQ ID NO:1815,SEQ ID NO:1816,SEQ ID NO:1817,SEQ ID NO:1818,SEQ ID NO:1819,SEQ ID NO:1820,SEQ ID NO:1821,SEQ ID NO:1822|RAF1,PKS2,RAFA1|12,36,104,166|
 SEQ ID NO:182|13|SEQ ID NO:1823,SEQ ID NO:1824,SEQ ID NO:1825,SEQ ID NO:1826,SEQ ID NO:1827,SEQ ID NO:1828,SEQ ID NO:1829,SEQ ID NO:1830,SEQ ID NO:1831,SEQ ID NO:1832,SEQ ID NO:1833,SEQ ID NO:1834,SEQ ID NO:1835|EIF3-P46,EIF3-P48,EIF3S6,INT6|72,231,336,391,491,
 492|
 SEQ ID NO:183|13|SEQ ID NO:1775,SEQ ID NO:1776,SEQ ID NO:1777,SEQ ID NO:1778,SEQ ID NO:1779,SEQ ID NO:1780,SEQ ID NO:1782,SEQ ID NO:1783,SEQ ID NO:1784|CTNNB,CTNNB1|1,2,8,10,
 16,36,89,151,154,186,790|
 10 SEQ ID NO:184|13|SEQ ID NO:1836,SEQ ID NO:1837,SEQ ID NO:1838,SEQ ID NO:1839,SEQ ID NO:1840,SEQ ID NO:1841,SEQ ID NO:1842|ERM,ETV5|3,7,23|
 SEQ ID NO:185|13|SEQ ID NO:1843,SEQ ID NO:1844,SEQ ID NO:1845,SEQ ID NO:1846,SEQ ID NO:1847,SEQ ID NO:1848,SEQ ID NO:1849,SEQ ID NO:1850,SEQ ID NO:1851,SEQ ID NO:1852,SEQ ID NO:1853,SEQ ID NO:1854|R1AA0619,P160ROCK,ROCK1,ROCK2|12,87,89,104,105,131,132,133,134,320,
 502,581,583,678|
 SEQ ID NO:186|13|SEQ ID NO:1855,SEQ ID NO:1856,SEQ ID NO:1857,SEQ ID NO:1858,SEQ ID NO:1859,SEQ ID NO:1860,SEQ ID NO:1861|PFN1|134,137,857|
 SEQ ID NO:187|14|SEQ ID NO:682,SEQ ID NO:683,SEQ ID NO:684,SEQ ID NO:685,SEQ ID NO:686,
 SEQ ID NO:1862,SEQ ID NO:1863,SEQ ID NO:1864,SEQ ID NO:687,SEQ ID NO:688|CD71,TFR,TFRC|31,
 331,590,720,721,722|
 15 SEQ ID NO:188|14|SEQ ID NO:929,SEQ ID NO:930,SEQ ID NO:931,SEQ ID NO:932,SEQ ID NO:933,
 SEQ ID NO:934,SEQ ID NO:935,SEQ ID NO:936,SEQ ID NO:937,SEQ ID NO:938,SEQ ID NO:939|CD49F,
 ITGA6|34,200,224,405,408,923|
 SEQ ID NO:189|15|SEQ ID NO:1865,SEQ ID NO:1866,SEQ ID NO:1867,SEQ ID NO:1868,SEQ ID NO:1869,SEQ ID NO:1870,SEQ ID NO:1871,SEQ ID NO:1872,SEQ ID NO:1873,SEQ ID NO:1874,SEQ ID NO:1875,SEQ ID NO:1876,SEQ ID NO:1877,SEQ ID NO:1878,SEQ ID NO:1879,SEQ ID NO:1880,SEQ ID NO:1881,SEQ ID NO:1882|GLBA,PSAP,SAP1|17,209,234,369,377,438,464,1181|
 SEQ ID NO:190|15|SEQ ID NO:1462,SEQ ID NO:1464,SEQ ID NO:1466,SEQ ID NO:1468,SEQ ID NO:1472,SEQ ID NO:1475,SEQ ID NO:1476,SEQ ID NO:1478,SEQ ID NO:1479,SEQ ID NO:1482,SEQ ID NO:1483,SEQ ID NO:1485,SEQ ID NO:1487,SEQ ID NO:1488,SEQ ID NO:1489,SEQ ID NO:1491,SEQ ID NO:1493,SEQ ID NO:1496,SEQ ID NO:1500,SEQ ID NO:1503|CD66E,CEA,CEACM5|31,297|
 SEQ ID NO:191|15|SEQ ID NO:1340,SEQ ID NO:1341,SEQ ID NO:1342,SEQ ID NO:1343,SEQ ID NO:1344,SEQ ID NO:1345,SEQ ID NO:1346,SEQ ID NO:1347,SEQ ID NO:1348,SEQ ID NO:1350,SEQ ID NO:1351,SEQ ID NO:1352,SEQ ID NO:1353|GALBP,LGALS2,LGALS3,MAC-2,MAC2|56,154,635,1012|
 20 SEQ ID NO:192|13|SEQ ID NO:1883,SEQ ID NO:1462,SEQ ID NO:1464,SEQ ID NO:1465,SEQ ID NO:1466,SEQ ID NO:1467,SEQ ID NO:1468,SEQ ID NO:1884,SEQ ID NO:1472,SEQ ID NO:1885,SEQ ID NO:1886,SEQ ID NO:1887,SEQ ID NO:1475,SEQ ID NO:1679,SEQ ID NO:1476,SEQ ID NO:1478,SEQ ID
 25
 30
 35

5 NO:1888, SEQ ID NO:1479, SEQ ID NO:1889, SEQ ID NO:1482, SEQ ID NO:1483, SEQ ID NO:1484, SEQ ID NO:1485, SEQ ID NO:1487, SEQ ID NO:1488, SEQ ID NO:1489, SEQ ID NO:1490, SEQ ID NO:1491, SEQ ID NO:1493, SEQ ID NO:1890, SEQ ID NO:1891, SEQ ID NO:1495, SEQ ID NO:1496, SEQ ID NO:1892, SEQ ID NO:1893, SEQ ID NO:1500, SEQ ID NO:1894, SEQ ID NO:1503, SEQ ID NO:1895, SEQ ID NO:1896, SEQ ID NO:1507|CD66E, CEA, CEACAM5, CEACAM7, CGM2|2,31,154,209,297|
 SEQ ID NO:193|15|SEQ ID NO:1340, SEQ ID NO:1341, SEQ ID NO:1342, SEQ ID NO:1343, SEQ ID NO:1344, SEQ ID NO:1345, SEQ ID NO:1346, SEQ ID NO:1347, SEQ ID NO:1348, SEQ ID NO:1350, SEQ ID NO:1344, SEQ ID NO:1351, SEQ ID NO:1352, SEQ ID NO:1353|GALBP, LGALS2, LGALS3, MAC-2, MAC2|56,
 154,635,1012|

10 SEQ ID NO:194|15|SEQ ID NO:929, SEQ ID NO:930, SEQ ID NO:931, SEQ ID NO:932, SEQ ID NO:933, SEQ ID NO:1774, SEQ ID NO:934, SEQ ID NO:935, SEQ ID NO:936, SEQ ID NO:937, SEQ ID NO:938, SEQ ID NO:939|CD49F, ITGA6|34,200,224,405,408,923|
 SEQ ID NO:195|15|SEQ ID NO:1897, SEQ ID NO:1898, SEQ ID NO:1899|ALCAM, CD166, MEMD|31,89,125,
 129,141,147,186,187,387|
 15 SEQ ID NO:196|15|SEQ ID NO:1242, SEQ ID NO:1462, SEQ ID NO:1463, SEQ ID NO:1464, SEQ ID NO:1465, SEQ ID NO:1466, SEQ ID NO:1676, SEQ ID NO:1243, SEQ ID NO:1470, SEQ ID NO:1900, SEQ ID NO:1472, SEQ ID NO:1901, SEQ ID NO:1475, SEQ ID NO:1679, SEQ ID NO:1476, SEQ ID NO:1477, SEQ ID NO:1478, SEQ ID NO:1480, SEQ ID NO:1482, SEQ ID NO:1244, SEQ ID NO:1483, SEQ ID NO:1902, SEQ ID NO:1485, SEQ ID NO:1486, SEQ ID NO:1487, SEQ ID NO:1903, SEQ ID NO:1488, SEQ ID NO:1489, SEQ ID NO:1491, SEQ ID NO:1492, SEQ ID NO:1493, SEQ ID NO:1496, SEQ ID NO:1497, SEQ ID NO:1500, SEQ ID NO:1502, SEQ ID NO:1503, SEQ ID NO:1505, SEQ ID NO:1700, SEQ ID NO:1506, SEQ ID NO:1904, SEQ ID NO:1245|CD66C, CD66E, CEA, CEACAM5, CEACAM6, NCA|31,34,89,120,297,389|
 SEQ ID NO:197|15|SEQ ID NO:1340, SEQ ID NO:1341, SEQ ID NO:1342, SEQ ID NO:1343, SEQ ID NO:1344, SEQ ID NO:1345, SEQ ID NO:1346, SEQ ID NO:1347, SEQ ID NO:1348, SEQ ID NO:1349, SEQ ID NO:1350, SEQ ID NO:1351, SEQ ID NO:1352, SEQ ID NO:1353|GALBP, LGALS2, LGALS3, MAC-2, MAC2|56,
 154,635,1012|
 SEQ ID NO:198|15|SEQ ID NO:1702, SEQ ID NO:1703, SEQ ID NO:1704, SEQ ID NO:1705, SEQ ID NO:1706, SEQ ID NO:1707, SEQ ID NO:1708, SEQ ID NO:1709, SEQ ID NO:1710, SEQ ID NO:1711, SEQ ID NO:1712, SEQ ID NO:1713|5F7, BSG, CD147,OK|89,242,386,739|
 20 SEQ ID NO:199|8|SEQ ID NO:1905, SEQ ID NO:1906, SEQ ID NO:1907, SEQ ID NO:1908, SEQ ID NO:1912|CD45, GP180,LCA,PTPRC,T200|31,34,47,
 386,469,858|
 SEQ ID NO:200|13|SEQ ID NO:1745, SEQ ID NO:1746, SEQ ID NO:1913, SEQ ID NO:1747, SEQ ID NO:1748, SEQ ID NO:1749, SEQ ID NO:1914, SEQ ID NO:1750, SEQ ID NO:1751, SEQ ID NO:1915, SEQ ID NO:1752, SEQ ID NO:1916, SEQ ID NO:1753|ADAM17, CD156B, CSVP, TACE|31,47,51,120,242,389,640|
 35 SEQ ID NO:201|8|SEQ ID NO:256, SEQ ID NO:257, SEQ ID NO:258, SEQ ID NO:259, SEQ ID NO:260, SEQ ID NO:261, SEQ ID NO:262, SEQ ID NO:263, SEQ ID NO:264|CD27L, CD27LG, CD70, KI-24 ANTIGEN,

TNFSF7 | 31, 56, 57, 89, 120, 179, 387, 388, 389 |
 SEQ ID NO:202 | 8 | SEQ ID NO:256, SEQ ID NO:257, SEQ ID NO:259, SEQ ID NO:260, SEQ
 ID NO:261, SEQ ID NO:262, SEQ ID NO:263, SEQ ID NO:264 | CD27L, CD27LG, CD70, KI-24 ANTIGEN,
 TNFSF7 | 31, 56, 57, 89, 120, 179, 387, 388, 389 |
 5 SEQ ID NO:203 | 8 | SEQ ID NO:1515, SEQ ID NO:1516, SEQ ID NO:1517, SEQ ID NO:1518, SEQ ID
 ID NO:890, SEQ ID NO:1520, SEQ ID NO:1521, SEQ ID NO:1522, SEQ ID NO:1523, SEQ ID NO:1524, SEQ ID
 NO:896, SEQ ID NO:897, SEQ ID NO:898, SEQ ID NO:899, SEQ ID NO:900, SEQ ID NO:901, SEQ ID
 NO:902, SEQ ID NO:903, SEQ ID NO:904 | CD23, CD23A, FCE2, FCER2, SST | 28, 31, 33, 34, 55, 56, 102, 120,
 129, 179, 253, 359, 386, 389, 438, 765, 848, 849, 850, 851 |
 15 SEQ ID NO:205 | 8 | SEQ ID NO:498, SEQ ID NO:499, SEQ ID NO:500, SEQ ID NO:501, SEQ ID NO:502, SEQ
 ID NO:503, SEQ ID NO:504, SEQ ID NO:505, SEQ ID NO:506, SEQ ID NO:507, SEQ ID NO:508, SEQ ID
 NO:509, SEQ ID NO:510 | 4F2, 4F2HC, CD98, MDU1, NACAE, SLC3A2 |
 SEQ ID NO:206 | 8 | SEQ ID NO:498, SEQ ID NO:499, SEQ ID NO:500, SEQ ID NO:501, SEQ ID NO:502, SEQ
 ID NO:503, SEQ ID NO:504, SEQ ID NO:1917, SEQ ID NO:1918, SEQ ID NO:505, SEQ ID NO:506, SEQ ID
 NO:507, SEQ ID NO:508, SEQ ID NO:509, SEQ ID NO:510 | 4F2, 4F2HC, CD98, MDU1, NACAE, SLC3A2 |
 20 SEQ ID NO:207 | 8 | SEQ ID NO:1515, SEQ ID NO:1516, SEQ ID NO:1517, SEQ ID NO:1518, SEQ ID
 NO:1519, SEQ ID NO:1919, SEQ ID NO:1920, SEQ ID NO:1520, SEQ ID NO:1521, SEQ ID NO:1522, SEQ ID
 NO:1523, SEQ ID NO:1921, SEQ ID NO:1524, SEQ ID NO:1525, SEQ ID NO:1526, SEQ ID NO:1527, SEQ ID
 NO:1528, SEQ ID NO:1529, SEQ ID NO:1530, SEQ ID NO:1531, SEQ ID NO:1533, SEQ ID NO:1534, SEQ ID
 NO:1535 | B1, BP35, CD20, MS4A1, MS4A1A, MS4A2 | 22, 31, 57, 129, 359, 789, 1110 |
 SEQ ID NO:208 | 8 | SEQ ID NO:1922, SEQ ID NO:1923, SEQ ID NO:1924, SEQ ID NO:1925, SEQ ID
 ID NO:1926, SEQ ID NO:1927, SEQ ID NO:1928, SEQ ID NO:1929, SEQ ID NO:1930, SEQ ID NO:1931, SEQ ID
 NO:1932 | 4F9, C33, CD82, GR15, IA4, KAI1, PREB, R2, SAR2, ST6 | 2, 8, 10, 14, 16, 17, 31, 129, 154 |
 SEQ ID NO:209 | 8 | SEQ ID NO:1933, SEQ ID NO:1934, SEQ ID NO:1935, SEQ ID NO:1936, SEQ ID
 30 NO:1937, SEQ ID NO:1938, SEQ ID NO:1939, SEQ ID NO:1940, SEQ ID NO:1941, SEQ ID NO:1942, SEQ ID
 NO:1943, SEQ ID NO:1944 | CD74, DHLAG | 129, 209, 413 |
 SEQ ID NO:210 | 8 | SEQ ID NO:498, SEQ ID NO:499, SEQ ID NO:500, SEQ ID NO:501, SEQ ID NO:503, SEQ
 ID NO:504, SEQ ID NO:505, SEQ ID NO:506, SEQ ID NO:507, SEQ ID NO:508, SEQ ID NO:510 | 4F2, 4F2HC,
 CD98, MDU1, NACAE, SLC3A2 |
 35 SEQ ID NO:211 | 8 | SEQ ID NO:708, SEQ ID NO:709, SEQ ID NO:710, SEQ ID NO:711, SEQ ID NO:712, SEQ
 ID NO:713, SEQ ID NO:714, SEQ ID NO:715, SEQ ID NO:716, SEQ ID NO:717, SEQ ID NO:718, SEQ ID
 NO:719, SEQ ID NO:720 | CD18, ITGB2, LAD, LOAMB, LFA-1, MF17 | 34, 129, 141, 186, 405, 408, 766 |

- 5 NO:747, SEQ ID NO:1994, SEQ ID NO:1995, SEQ ID NO:1996, SEQ ID NO:1997, SEQ ID NO:1998, SEQ ID NO:1999, SEQ ID NO:2000, SEQ ID NO:2001, SEQ ID NO:748, SEQ ID NO:749, SEQ ID NO:750, SEQ ID NO:751, SEQ ID NO:752, SEQ ID NO:2002, SEQ ID NO:754 |BB2, CD54, ICAM1|129,154,186,187,219,231,331,385,717,778| SEQ ID NO:222|8|SEQ ID NO:2003, SEQ ID NO:2004, SEQ ID NO:2005, SEQ ID NO:2006, SEQ ID NO:2007, SEQ ID NO:2008, SEQ ID NO:2009, SEQ ID NO:2010, SEQ ID NO:2011|CD81,TAPA1|31,56,57,129,154,192,332|
- 10 SEQ ID NO:223|8|SEQ ID NO:498, SEQ ID NO:499, SEQ ID NO:500, SEQ ID NO:503, SEQ ID NO:504, SEQ ID NO:505, SEQ ID NO:506, SEQ ID NO:508, SEQ ID NO:510|4F2, 4F2HC, CD98, MDU1,NACAE, SLC3A2| SEQ ID NO:224|8|SEQ ID NO:1905, SEQ ID NO:1906, SEQ ID NO:1907, SEQ ID NO:1908, SEQ ID NO:1909, SEQ ID NO:1910, SEQ ID NO:1911, SEQ ID NO:1912|CD45, GP180,LCA, PTPRC, T200|31,34,47,386,469,858|
- 15 SEQ ID NO:225|8|SEQ ID NO:498, SEQ ID NO:499, SEQ ID NO:500, SEQ ID NO:501, SEQ ID NO:502, SEQ ID NO:503, SEQ ID NO:504, SEQ ID NO:505, SEQ ID NO:506, SEQ ID NO:508, SEQ ID NO:509, SEQ ID NO:510|4F2, 4F2HC, CD98, MDU1,NACAE, SLC3A2|
- 20 SEQ ID NO:226|8|SEQ ID NO:498, SEQ ID NO:499, SEQ ID NO:500, SEQ ID NO:501, SEQ ID NO:502, SEQ ID NO:503, SEQ ID NO:504, SEQ ID NO:505, SEQ ID NO:506, SEQ ID NO:507, SEQ ID NO:508, SEQ ID NO:509, SEQ ID NO:510|4F2, 4F2HC, CD98, MDU1,NACAE, SLC3A2|
- 25 SEQ ID NO:227|8|SEQ ID NO:256, SEQ ID NO:258, SEQ ID NO:262, SEQ ID NO:264|CD27L,CD27LG, CD70, KI-24 ANTIGEN, TNFSF7|31,56,57,89,120,179,387,388,389| SEQ ID NO:228|8|SEQ ID NO:2012, SEQ ID NO:2013, SEQ ID NO:2014, SEQ ID NO:2015, SEQ ID NO:2016, SEQ ID NO:2017, SEQ ID NO:2018, SEQ ID NO:2019, SEQ ID NO:2020, SEQ ID NO:2021, SEQ ID NO:2022, SEQ ID NO:2023, SEQ ID NO:2024, SEQ ID NO:2025, SEQ ID NO:2026, SEQ ID NO:2027, SEQ ID NO:2028, SEQ ID NO:2029, SEQ ID NO:2030, SEQ ID NO:2031|CD22, SIGLEC-2|31,120,129,141,186,187|
- 30 SEQ ID NO:229|8|SEQ ID NO:1933, SEQ ID NO:1934, SEQ ID NO:1935, SEQ ID NO:1936, SEQ ID NO:1937, SEQ ID NO:1938, SEQ ID NO:1939, SEQ ID NO:2032, SEQ ID NO:1940, SEQ ID NO:1941, SEQ ID NO:1942, SEQ ID NO:1943, SEQ ID NO:1944|CD74,DHLAG|129,209,413| SEQ ID NO:230|8|SEQ ID NO:1735, SEQ ID NO:1736, SEQ ID NO:1737, SEQ ID NO:1738, SEQ ID NO:1739, SEQ ID NO:1740, SEQ ID NO:1741, SEQ ID NO:2033, SEQ ID NO:1742, SEQ ID NO:1743, SEQ ID NO:1744|CD11A, ITGAL, LFA-1, LFA1A|34,161,186,201,405,408|
- 35 SEQ ID NO:231|8|SEQ ID NO:1905, SEQ ID NO:1906, SEQ ID NO:1907, SEQ ID NO:1910, SEQ ID NO:1911, SEQ ID NO:2035, SEQ ID NO:1908, SEQ ID NO:2036, SEQ ID NO:1909, SEQ ID NO:1912|CD45, GP180,LCA,PTPRC, T200|31,34,47,386,469,858|
- SEQ ID NO:232|8|SEQ ID NO:498, SEQ ID NO:499, SEQ ID NO:500, SEQ ID NO:501, SEQ ID NO:502, SEQ ID NO:503, SEQ ID NO:504, SEQ ID NO:505, SEQ ID NO:506, SEQ ID NO:507, SEQ ID NO:508, SEQ ID NO:509, SEQ ID NO:510|4F2, 4F2HC, CD98, MDU1,NACAE, SLC3A2|
- SEQ ID NO:233|8|SEQ ID NO:1268, SEQ ID NO:1269, SEQ ID NO:1270, SEQ ID NO:1271, SEQ ID

NO:1272, SEQ ID NO:1273, SEQ ID NO:1274, SEQ ID NO:1275, SEQ ID NO:1276, SEQ ID NO:1277, SEQ ID NO:1278, SEQ ID NO:1279, SEQ ID NO:1280, SEQ ID NO:2037, SEQ ID NO:1281, SEQ ID NO:1282, SEQ ID NO:1283, SEQ ID NO:1284, SEQ ID NO:1285, SEQ ID NO:1286, SEQ ID NO:1287, SEQ ID NO:1288, SEQ ID NO:1289, SEQ ID NO:1290, SEQ ID NO:1291, SEQ ID NO:1292, SEQ ID NO:1293, SEQ ID NO:2039, SEQ ID NO:1296, SEQ ID NO:1297, SEQ ID NO:1298, SEQ ID NO:1299, SEQ ID NO:1300, SEQ ID NO:1301, SEQ ID NO:1302, SEQ ID NO:1303, SEQ ID NO:1304, SEQ ID NO:1305, SEQ ID NO:1306, SEQ ID NO:1307, SEQ ID NO:1308, SEQ ID NO:1309, SEQ ID NO:1310, SEQ ID NO:1311, SEQ ID NO:2040, SEQ ID NO:1312, SEQ ID NO:1313, SEQ ID NO:1314, SEQ ID NO:1315, SEQ ID NO:1316, SEQ ID NO:1317, SEQ ID NO:1318, SEQ ID NO:1319|CD44, CD44R, IN, MC56, MDU2, MDU3, MTC4|
SEQ ID NO:234|8|SEQ ID NO:2041|GLBA, PSAP, SAP1|17, 209, 234, 369, 377, 438, 464, 1181|
SEQ ID NO:235|18|SEQ ID NO:940, SEQ ID NO:943, SEQ ID NO:945, SEQ ID NO:948, SEQ ID NO:949, SEQ ID NO:950, SEQ ID NO:951, SEQ ID NO:958, SEQ ID NO:959, SEQ ID NO:960, SEQ ID NO:961, SEQ ID NO:962, SEQ ID NO:963, SEQ ID NO:964, SEQ ID NO:965, SEQ ID NO:970, SEQ ID NO:971, SEQ ID NO:972|CALM, CLTH,
10 15 DJ167A19.1, FLT4, KIAA0656, PCL, PICALM, SNAP91, VEGFR3 | 12, 31, 34, 135, 332, 411, 461, 639, 815 |

Other Embodiments

While the invention has been described in conjunction with the detailed description thereof, the foregoing description is intended to illustrate and not limit the scope of the invention, which is defined by the scope of the appended claims. Other aspects, advantages, and modifications are within the scope of the following claims.

5

What is claimed is:

1. An isolated nucleic acid comprising a nucleotide sequence that encodes a polypeptide comprising an amino acid sequence which is at least 95% identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.
- 5 2. An isolated nucleic acid comprising a nucleotide sequence that encodes a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.
- 10 3. The nucleic acid of claim 2, wherein the peptide sequence is identical to that of a naturally processed class I or class II MHC-binding peptide.
- 15 4. An isolated nucleic acid comprising a nucleotide sequence that encodes a polypeptide comprising an amino acid sequence selected from the group consisting of a variant of any one of SEQ ID NOs: 1-235, wherein the variant has no more than two conservative amino acid substitutions.
- 20 5. An isolated nucleic acid comprising a nucleotide sequence that encodes a polypeptide comprising at least 8 contiguous amino acids of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.
- 25 6. An isolated nucleic acid comprising a nucleotide sequence encoding a polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.
- 30 7. An isolated nucleic acid comprising a nucleotide sequence encoding a polypeptide consisting essentially of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.
8. An isolated nucleic acid comprising a nucleotide sequence encoding a polypeptide comprising no more than 30 contiguous amino acids of a naturally occurring

human protein, wherein the naturally occurring protein comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

9. The nucleic acid of claim 8, wherein the peptide sequence is identical to that of
5 a naturally processed class I or class II MHC-binding peptide.

10. A purified polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

10 11. The polypeptide of claim 10, wherein the peptide sequence is identical to that of a naturally processed class I or class II MHC-binding peptide.

12. A purified polypeptide comprising at least 8 contiguous amino acids of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.

15 13. A purified polypeptide, comprising at least an immunogenic portion of a protein, wherein the protein comprises an amino acid selected from the group consisting of SEQ ID NOs: 1-235.

20 14. A purified immunogenic polypeptide comprising at least 8 contiguous amino acids of an amino acid sequence selected from the group consisting of SEQ ID NOS: 1-235.

25 15. A purified polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235, wherein the purified polypeptide comprises at least 25 amino acids.

16. The purified polypeptide of claim 14 wherein the polypeptide comprises fewer than 100 amino acids.

30

17. The purified polypeptide of claim 15 wherein the polypeptide comprises fewer than 50 amino acids.
18. A purified polypeptide consisting of an amino acid sequence selected from 5 the group consisting of SEQ ID NOs:1-235.
19. A purified polypeptide consisting essentially of an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235.
- 10 20. A vector comprising the nucleic acid of claim 1.
21. The vector of claim 20 wherein the vector comprises expression control sequences that direct the expression of the polypeptide.
- 15 22. The vector of claim 20 wherein the vector comprises expression control sequences that direct expression of the nucleic acid molecule.
23. A cell comprising the vector of claim 20.
- 20 24. An antibody that selectively binds a polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235.
- 25 25. A method of making an antibody, the method comprising:
 - (a) providing a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235 to a mammal in an amount effective to induce the production of an antibody that binds to the polypeptide;
 - (b) isolating from the mammal a cell that produces an antibody that selectively binds to a polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235;
 - 30 (c) immortalizing the cell isolated in step (b); and
 - (d) isolating antibodies from the immortalized cell.

26. The antibody of claim 24 wherein the polypeptide is expressed on a cell surface.
27. The antibody of claim 24, wherein the polypeptide is a target of a second
5 antibody located on a cell surface.
28. A humanized antibody which specifically binds to a domain of a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-
235.
10
29. The humanized antibody of claim 28 which is a full length antibody.
30. The humanized antibody of claim 28 which is a human IgG.
15
31. The humanized antibody of claim 28 which is an antibody fragment.
32. The humanized antibody of claim 28 wherein the antibody fragment is a
F(ab')₂.
- 20
33. A labeled antibody comprising the humanized antibody of claim 28 bound to a detectable label.
34. An immobilized antibody comprising the humanized antibody of claim 28
bound to a solid phase.
25
35. A conjugate comprising the humanized antibody of claim 28 bound to a cytotoxic agent.
36. A method for determining the presence of a protein comprising exposing a
30 sample suspected of containing the protein to the humanized antibody of claim 28 and determining binding of said antibody to the sample.

37. A kit comprising the humanized antibody of claim 28 and instructions for using the humanized antibody to detect a protein that binds to the antibody.

5 38. An isolated nucleic acid encoding the humanized antibody of claim 28.

39. A method for modulating the activity of the polypeptide of claim 10, the method comprising contacting the polypeptide with a compound that binds to the polypeptide in a concentration sufficient to modulate the activity of the polypeptide.

10

40. The method of claim 39, wherein the compound that binds the polypeptide is an antibody that selectively binds a polypeptide consisting of an amino acid sequence selected for the group consisting of SEQ ID NOs:1-235.

15

41. A method of treating a disorder in a mammal, the method comprising:

(a) identifying a mammal with the disorder; and

(b) administering to the mammal a compound that modulates the expression or activity of the polypeptide of claim 10,

wherein the administration results in an amelioration of one or more symptoms of
20 the disorder.

42. A method for detecting the presence of a polypeptide of claim 10 in a sample, the method comprising:

25 contacting the sample with a compound that selectively binds to a polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235; and

determining whether the compound binds to the polypeptide in the sample.

30 43. A method for detecting the presence of a disorder in a mammal, the method comprising:

(a) providing a biological sample derived from the mammal;

- (b) contacting the sample with a compound that binds to the polypeptide of claim 17 or to a nucleic acid that encodes the polypeptide of claim 17; and
(c) determining whether the compound binds to the sample,
wherein binding of the compound to the sample indicates the presence or absence of 5 the disorder in the mammal.

44. A method for imaging a site in a mammal, the method comprising:
(a) administering a compound to a mammal, wherein the compound binds to the polypeptide of claim 10 or to a nucleic acid that encodes the polypeptide; and
10 (b) detecting the compound with an imaging detector, thereby imaging the site in the mammal.

45. A method for identifying a compound that modulates the activity of the polypeptide of claim 10, the method comprising:
15 (a) contacting the polypeptide of claim 8 with a test compound; and
(b) determining the effect of the test compound on the activity of the polypeptide, to thereby identify a compound that modulates the activity of the polypeptide.

46. A method for identifying a compound that modulates the expression of the 20 nucleic acid of claim 2, the method comprising:
contacting the nucleic acid of claim 2 with a test compound; and
determining the effect of the test compound on the expression of the nucleic acid, to thereby identify a compound that modulates the expression of the nucleic acid.

- 25 47. A polypeptide profile that is characteristic of a given cell, wherein the profile comprises a representation of at least ten different polypeptides in the cell, wherein each of the at least ten different polypeptides comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235, and wherein the polypeptide profile is a reproducible characteristic of the cell.

30

48. The polypeptide profile of claim 47, wherein the each of the at least ten different polypeptides comprises an MHC-binding polypeptide.

49. The polypeptide profile of claim 47, wherein the representation characterizes 5 each individual polypeptide based upon at least one physical or chemical attribute of the polypeptide, the at least one physical or chemical attribute comprising an amino acid sequence.

50. The polypeptide profile of claim 47, wherein the representation characterizes 10 each individual peptide based upon at least two physical or chemical attributes.

51. The polypeptide profile of claim 47, wherein one of the physical or chemical attributes is a nucleotide sequence encoding the amino acid sequence.

15 52. The polypeptide profile of claim 47, wherein one of the physical or chemical attributes is mass-to-charge ratio.

53. The polypeptide profile of claim 47, wherein one of the physical or chemical attributes is an ion-fragmentation pattern.

20 54. The polypeptide profile of claim 47, wherein the representation characterizes each individual peptide based upon at least three physical or chemical attributes.

55. A database, stored on a machine-readable medium, comprising:
25 (a) two categories of data respectively representing: (i) peptide profiles and (ii) cell sources; and
(b) associations among instances of the two categories of data,
wherein the data representing polypeptide profiles comprise the peptide profile of
claim 46, and
30 wherein the database configures a computer to enable finding instances of data of
one of the categories based on their associations with instances of data the other category.

56. A method of selecting an antibody, the method comprising:
(a) contacting the polypeptide of claim 10 with an in vitro library of antibodies;
(b) binding an antibody to the polypeptide; and
5 (c) selecting the antibody that binds to the polypeptide.
57. An immunogenic composition comprising a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235, the composition when injected into a mammal eliciting an immunogenic response directed against a
10 polypeptide consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-235.
58. A method for treating a cancer comprising administering to a patient an amount of a composition comprising a polypeptide comprising an amino acid sequence
15 selected from the group consisting of SEQ ID NOs:1-235 in an amount sufficient to elicit an immunogenic response.
59. A method for treating a cancer patient, the method comprising administering to the patient an antibody that selectively binds to a peptide consisting of an amino acid
20 sequence selected from the group consisting of SEQ ID NOs:1-235.
60. A peptide array comprising at least 100 peptides selected from the group consisting of peptides consisting of an amino acid sequence selected from the group
25 consisting of SEQ ID NOs:1-235, each peptide linked to a solid support at a known location.
61. A collection of at least 10 polypeptide arrays, each array comprising at least 100 polypeptides consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235, each peptide linked to a solid support at a known location.
30

62. A method for identifying a compound that binds to a naturally processed class I MHC-binding polypeptide, the method comprising exposing a test compound to a collection of at least 100 polypeptides selected from the group consisting of polypeptides having an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235, 5 and identifying a peptide to which the test compound binds.

63. A method for identifying a compound that binds to a naturally processed class II MHC-binding polypeptide, the method comprising exposing a test compound to a collection of at least 100 polypeptides selected from the group consisting of polypeptides 10 having an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235 and identifying a peptide to which the test compound binds.

64. A database, stored on a machine-readable medium, comprising:
(a) three categories of data respectively representing (i) polypeptides, (ii) cell 15 sources, and (iii) cell treatments; and
(b) associations among instances of the three categories of data,
wherein the data representing peptides comprises at least 100 polypeptides each having an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235, and
20 wherein the database configures a computer to enable finding instances of data of one of the categories based on their associations with instances of data of at least one other category.

65. A polypeptide profile that is characteristic of a selected cell under selected 25 conditions, wherein the profile comprises a representation of at least ten different polypeptides expressed by the cell, wherein each of the at least ten different polypeptides comprising an amino acid sequence selected from the group consisting of SEQ ID NOs:1-235, and wherein the polypeptide profile is a reproducible characteristic of the cell.

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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.



WO 2002/078524 A3

(54) Title: TRANSLATIONAL PROFILING

(57) Abstract: Polypeptides representative of proteins expressed by a given cell type and isolated nucleic acids that encode the polypeptides are disclosed. The compositions and method described can be used to define a cell type at a given developmental, metabolic, or disease stage by identifying and cataloging proteins expressed in the cell. The compositions can also be used in the manufacture of therapeutics as well as in diagnostics and drug screening.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US02/09671

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C07H 21/04; C07K 16/24
US CL : 536/22.1; 530/387.1

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
U.S. : 536/22.1; 530/387.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
Please See Continuation Sheet

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,969,110 A (BECKMANN et al.) 19 October 1999, see entire document.	1-9, 20-23
Y	US 6,312,912 B1 (HOPE et al) 06 November 2001, see entire document.	1-9, 20-23
Y	US 5,738,844 A (BECKMANN et al) 14 April 1998, see entire document.	1-9, 20-23
Y	US 6,124,528 A (SHEWMAKER) 26 September 2000, see entire document.	1-9, 20-23
Y, P	US 6,469,230 B1 (EDWARDS et al) 22 October 2002, see entire document	1-9, 20-23

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier application or patent published on or after the international filing date	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&"	document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means		
"P" document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

01 April 2004 (01.04.2004)

Date of mailing of the international search report

13 SEP 2004

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/US02/09671

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claim Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. Claim Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. Claim Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)This International Searching Authority found multiple inventions in this international application, as follows:
Please See Continuation Sheet

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 1-9, 20-23, in part

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

PCT/US02/09671

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Group I, claim(s) 1-9, 20-23, drawn to nucleic acids.

Group II, claim(s) 38, drawn to nucleic acids encoding 235 antibodies.

Group III, claim(s) 10-19, 24, 60, 61, 65, drawn to 235 polypeptides.

Group IV, claim(s) 26-35, 37, 57, drawn to 235 antibodies.

Group V, claim(s) 25, drawn to method of making 235 antibodies.

Group VI, claim(s) 36, 39, 40, 42-44, drawn to a method for determining the presence of 235 proteins.

Group VII, claim(s) 41, 58, 59, drawn to a method of treating a disorder using 235 antibodies.

Group VIII, claim(s) 45, 46, 56, 62, 63, drawn to a method for identifying a compound that modulates the activity of 235 polypeptides.

Group IX, claim(s) 55 and 64, drawn to a database.

This application contains claims directed to more than one species of the generic invention. These species are deemed to lack Unity of Invention because they are not so linked as to form a single inventive concept under PCT Rule 13.1. In order for more than one species to be searched, the appropriate additional search fees must be paid. The species are as follows:
The 235 different nucleic acids, proteins, antibodies.

The inventions listed as Groups I-IX do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: Beckman et al. (U>S> Patent 5,969,110)(October 19, 1999) teach an isolated nucleic acid comprising a nucleotide sequence that encodes a polypeptides comprising an amino acid sequence which is at least 95% identical to a given amino acid sequence as recited in Claim 1. Therefore, all other claims lack the same or corresponding special technical feature.

The species listed above do not relate to a single inventive concept under PCT Rule 13.1 because under PCT Rule 13.2, the species lack the same corresponding special technical features for the following reasons: Nucleic acids comprising a nucleotide sequence encoding a polypeptide comprising no more than 30 contiguous amino acids of a naturally occurring human protein has been disclosed by Beckmann et al.

Continuation of B. FIELDS SEARCHED Item 3:

Form PCT/ISA/210 (second sheet) (July 1998)

INTERNATIONAL SEARCH REPORT

PCT/US02/09671

WEST, STN, MEDLINE, BIOSIS, CAPLUS, EMBASE, GENBANK
nucleic acid, polypeptide, database, antibody

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